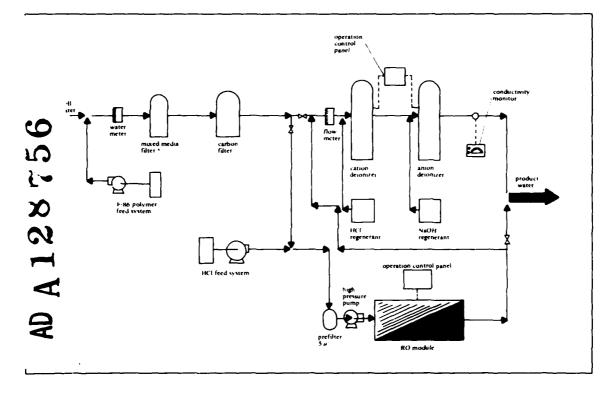


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Potable Water Treatment for Diego Garcia

by D. B. Chan and J. S. Williams January 1983



Technical Report R-898

Naval Civil Engineering Laboratory

Port Hueneme, California 93043

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INTRODUCTION

The water supply at Diego Garcia and at many other Naval installations in CONUS and overseas is very hard and has poor palatability due to contaminants, such as total dissolved solids (TDS), turbidity, color, taste, and odor. Hard water causes scaling in pipelines, reduces the life of boilers, cooling towers, and other heat exchange elements. On the other hand, high TDS, turbidity, color, taste, and odor reduce water palatability and affect personnel morale.

The wells supplying water at Diego Garcia possess all the contaminants described above and some exceed the U.S. Environmental Protection Agency's (EPA) and the Navy Bureau of Medicine (BUMED) recommended secondary drinking water standards. In addition, excessive quantities

of organics and microorganisms also exist in the raw well water.

The treated water quality now existing at Diego Garcia, however, does meet the primary (mandatory) drinking water standards and most of the recommended secondary drinking water standards. For the success of the Navy's mission and Fleet readiness, a high quality water that meets all of EPA secondary drinking water standards is desirable at Diego

Garcia if it can be feasibly attained.

The objective of this engineering investigation, sponsored by Pacific Division, Naval Facilities Engineering Command (PACNAVFACENGCOM) (Ref 1), is to develop a cost-effective water treatment system that would effectively remove or reduce TDS, color, taste, odor, turbidity, and the excessive amounts of organics and microorganisms so that all EPA standards can be met. The developed water treatment system will also have to meet other critical criteria: (a) minimum logistic support requirement, (b) low level skill and manpower to operate and maintain the system, and (c) capable of treating water containing high concentrations of TDS and other contaminants due to anticipated population increase and overdraft of wells causing water quality to degrade.

A series of water treatment components was evaluated by the Naval Civil Engineering Laboratory (NCEL), and a three-stage process was selected for test and evaluation at Diego Garcia. The system, consisting of a multi-media filter with a chemical feeder; a carbon filter and the demineralizing component, a cation-anion deionizer and a reverse osmosis (RO) system was fabricated. The water treatment system had a

design capacity of 20-gpm.

The water treatment system was installed at the Final Water Plant, Diego Garcia, on 13 March 1981, and was tested and evaluated during the period of 1 April and 5 August 1981. The water meters recorded that 570,000 gallons of water was produced by the deionizer and 1,033,000 gallons of water was produced by the RO system during the testing period. The product water possessed superior quality and far exceeded the EPA's standards. The results, conclusions, and recommendations of the water treatment system at Diego Garcia are summarized in this report.

BACKGROUND

The Navy's mission at Diego Garcia has expanded significantly during the last few years. This ever increasing military operation demands a larger quantity and a standard quality water supply for the personnel deployed on the island. At present the raw groundwater supply, which fulfills the larger quantity requirement, does not meet

all of EPA's secondary drinking water standards.

Some simple treatment systems are currently being used or are under construction at Diego Garcia. These systems include ultraviolet disinfection at the well fields, and aeration and chlorination at the Final Water Plant. The aeration and chlorination systems are currently in operation. The ultraviolet disinfection has been installed in most areas and should be in operation shortly. These systems treating and producing a composite water supply on Diego Garcia are expected to meet all secondary drinking water standards except those for removing TDS. It should be noted, however, that the aquifer has been degraded by locally high pumping rates and periodic droughts, causing a number of wells to upcone.

The groundwater quality at Diego Garcia is compared to EPA's

standards as follows:

| Parameters | EPA Standards | Diego Garcia Wells (Range) | Composite ^a of Wells (Range) |
|-----------------------------------|---------------------------------|----------------------------------|---|
| TDS (mg/l) | 500 | 270-1540 | 400-600 |
| Turbidity (JTV) | 1 | 3 | 1-3 |
| Color | 15 | +20 | 20-30 |
| 0dor | 3 | 3 [| 3 |
| Chloride (mg/l) | 250 | 25-400 ^D | 75-130 |
| Total Hardness (mg/l) | No Std. 150-300 is Hardwater | 160-480 | 160-200 |
| Microorganisms ^C (SPC) | No Std. | 10 ⁴ -10 ⁵ | |
| Total Coliform (MEC) | l coliform/100 ml | | <1 |
| Organic Chemicals ^C | No Std. | | |

^aWater sample taken from the raw water tank at the Final Water Plant.

The three-stage treatment process for this project was designed to function as follows:

bWell pumping instruction requires that pumping be terminated when chloride concentration exceeds 250 mg/l.

^CAdditional contaminants that need to be removed. Total hardness is directly related to the TDS and turbidity problem, while organic chemicals are the major cause of color, taste, odor and excessive growth of microorganisms.

| <u>Stage</u> | <u>Unit Process</u> a | Major Removal/Reduction Parameters |
|--------------|--|---|
| 1 | Multi-Media Filter with Chemical Feeder | Turbidity, microorganisms |
| 2 | Carbon Filter | Color, taste, odor, and organics |
| 3A | RO System | TDS, hardness, chloride, organics, microorganisms, turbidity, color, taste, and odor. |
| 3B | Cation-Anion Deionizer | TDS, hardness, chloride |

^aStage 1 and 2 unit processes are considered to be pretreatments for the RO and deionizer systems.

The RO process was selected for test and evaluation based on criteria established by NCEL. The RO possesses many advantages over other water purification methods; e.g., low energy consumption, low maintenance, is not sensitive to TDS concentration, etc. The RO like other membrane processes may require pretreatment of the feed water to prevent the membrane from fouling. Therefore, additional treatments (including pretreatment requirements) membrane fouling characteristics, and membrane cleaning requirements had to be evaluated prior to being used at Diego Garcia.

The ion-exchange deionization process was selected for testing because of its better known, field-proven technology. However, the ion exchange process requires a large quantity of chemical regenerants that could create a logistic burden on the Navy. The ion-exchange process may be used in the future to furnish high quality water for such uses as: injection water for aircraft takeoff, batteries, as an engine coolant, and boiler feed.

As indicated in the comparison of water quality between the wells (current raw water supplied at Diego Garcia) and EPA standards, the primary objective of this project is to reduce the TDS content in the water. In addition, the pretreatment required for the use of an ion-exchange deionizer or RO demineralizer would aid in the final removal of turbidity, color, taste, odor, and excessive organics. More importantly, should the groundwater quality degrade further due to upconing, drought, or man-made pollution, the three-stage water treatment system tested will provide the treatment required.

Specific objectives for the test and evaluation of the water treatment system are:

- 1. Determine the capability of the three-stage water treatment system to meet EPA/BUMED drinking water standards.
- 2. Identify additional treatments including pretreatment requirements.
- 3. Define ion-exchange resin and RO membrane fouling characteristics and their regeneration and or membrane cleaning requirements.

- 4. Determine reliability and maintainability of the water treatment system components and evaluate their applicability at Diego Garcia.
 - 5. Assess product water scaling and corrosion potential.
- 6. Analyze test data for preparation of design criteria for a feasible water treatment system.
- 7. Investigate* the total amount of trihalomethanes (TTHMs) in the well water, the water supply, and the product water of the water treatment system (before and after chlorination). The capability of the water treatment system to remove TTHMs and TTHMs' precursors must be determined.

TEST EQUIPMENT AND OPERATION**

Water Treatment System

Multi-Media Filter. The Culligan HD-20 Depth Filter, the tank on the right-hand side of Figure 1, has a design flow rate of 20 gpm, a loading rate of 10 gpm/ft², and a backwash rate of 30 gpm. The filter contains about 6 ft³ of multi-media materials, and the bed is approximately 33 inches deep. The filter tank is 20 inches in diameter and 54 inches long excluding the height of the supporting legs. An automated control system panel is built in for the filter operation and backwash. A space of 20 inches wide by 36 inches deep by 66 inches high is required for the filter. The filter can stand up to $100 \, \text{psi}$ operational pressure and temperatures ranging from $40 \, \text{°F}$ to $120 \, \text{°F}$. The saturated filter weighs $1,600 \, \text{pounds}$.

Activated Carbon Adsorber (Filter). The Culligan HR-24 Carbon Filter, the left hand-side tank of Figure 1, has a design flow rate of 20 gpm, a loading rate of 7 gpm/ft 2 , and a backwash rate of 30 gpm. The filter contains about 6.5 ft 3 of Cullar D carbon, and the bed is approximately 25 inches deep. The filter tank is 24-inches in diameter and 54 inches long excluding the supporting legs. An automated control system panel is built in for the filter operation and backwash. A space of 25 inches wide by 40 inches deep by 67 inches high is required for the filter. The filter can stand up to 100 psi operational pressure and temperatures ranging from $40^{\circ}\mathrm{F}$ to $120^{\circ}\mathrm{F}$. The saturated filter weighs about 2,150 pounds.

<u>Ion Exchange Deionizer</u>. Two Culligan AC-20W deionizers, Figure 2, are hooked up in a series, one for cation and the other for anion removal. The deionizers have a design flow rate of 10 gpm. The cation

^{*}This objective was added by Reference 2.

^{**}Further described in Reference 3.

resin (right-hand side tank) has an exchange capacity of about 230,000 grains expressed as $CaCO_3$, and the anion resin (left-hand side tank) has a capacity of 64,100 grains expressed as $CaCO_3$ per each regeneration cycle. The cation deionizer has 10 ft³ of exchange material (CH-1 resin), and requires 20 gallons of $20^{\circ}Be$ (31.4%) HCl to regenerate. The anion deionizer has 8 ft³ of exchange material (CW-4 resin), and requires 5 gallons of 50% NaOH to regenerate. Each of the resin tanks is 20 inches in diameter and 84 inches long excluding the supporting legs. An automated control system panel is attached for the deionizer operation and regeneration. A space of 78 inches wide by 32 inches deep by 102 inches high is required for the two deionizer resin tanks hooked up in a series. The deionizer can stand 100 psi operational pressure, and will tolerate a temperature range of $45^{\circ}F$ to $100^{\circ}F$. The deionizers weigh about 3,500 pounds each in operation.



Figure 1. Multi-media filter (right) and activated carbon adsorber (left).

Reverse Osmosis System. The Culligan KD-20 RO unit, Figure 3, has a design flow rate of 18 to 20 gpm. Nominal product water recovery is 75% initially. After 3 years, the recovery will be approximately 60%, due to membrane compaction and fouling. Final product water will contain about 10% of TDS in the feed water. The RO unit, a two-stage process, contains one DuPont B-9 hollow-fiber module (9-1/2 inches in diameter by 48 inches long by 47 inches wide) and one DuPont B-9 hollow-fiber module (5-1/4 inches in diameter by 47 inches long). The two stages are used to produce the maximum amount of product water from a given amount of feed water (one pass). The RO system is also equipped

with a 5 μ prefilter (cartridge) and a 15 hp motor-pump assembly. The RO unit can tolerate 400 psi operational pressure and temperatures ranging from 40°F to 95°F. A space of 120 inches by 36 inches deep by 80 inches high is required for installation of the RO unit. The unit weighs about 1,725 pounds (dry).

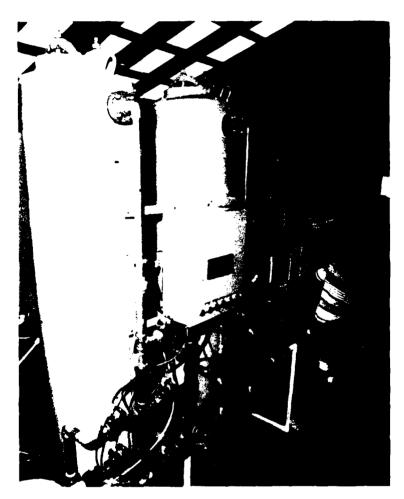


Figure 2. Ton exchange deionizer water treatment system

Accessories

Chemical Feeders and Feeder Systems. The Culligan DT-60A chemical feeder system was procured to feed F-86 into the feed water before R0 treatment. F-86 is a highly active cationic liquid polyelectrolyte used in conjunction with the multi-media filter to remove a majority of the colloidal particles. The feeder system has a capacity of 60 gal/day and is equipped with a 40-gal chemical storage tank. A feed rate determination test procedure was provided by Culligan. A total of 90 gallons of F-86 concentrate was procured for field testing at Diego Garcia.

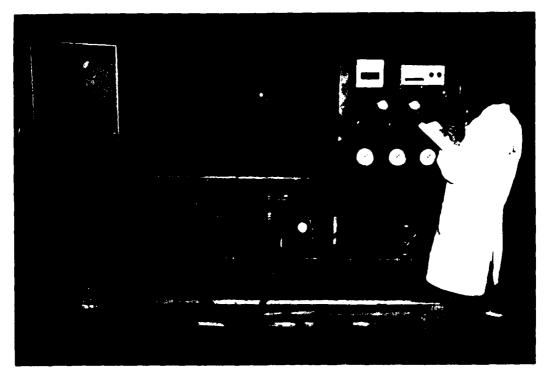


Figure 3. The reverse osmosis water treatment system.

Culligan Special pH Control. The Culligan pH monitor/controller was procured for use. The controller, with a shutdown feature, is designed to continuously feed 20° Be (31.4%) HCl at a preset rate, into the feed water of the RO unit. It will stop the acid feeder when the pH drops below a preset range of 5.2 to 5.6. The acid is used to control scale formation on the RO membrane.

Fouling Index Test Kit. Culligan supplied a fouling index test kit for determinating the quality of the feed water necessary to protect the RO membrane. Instructions for operating the test kit were provided by Culligan along with an RO unit operation manual.

Water Treatment System Installation and Operation

Installation and operation manuals for the water treatment system were provided by Culligan, and a list of these manuals can be found in Appendix A (a schematic of the water treatment system tested at Diego Garcia is shown in Figure 4).

TEST PROCEDURES, METHODOLOGY, AND RESULTS

Test and evaluation of the water treatment system at Diego Garcia was divided into six major parts. The test procedures, methodology, and sample collection and analysis are listed below and described in subsequent sections:

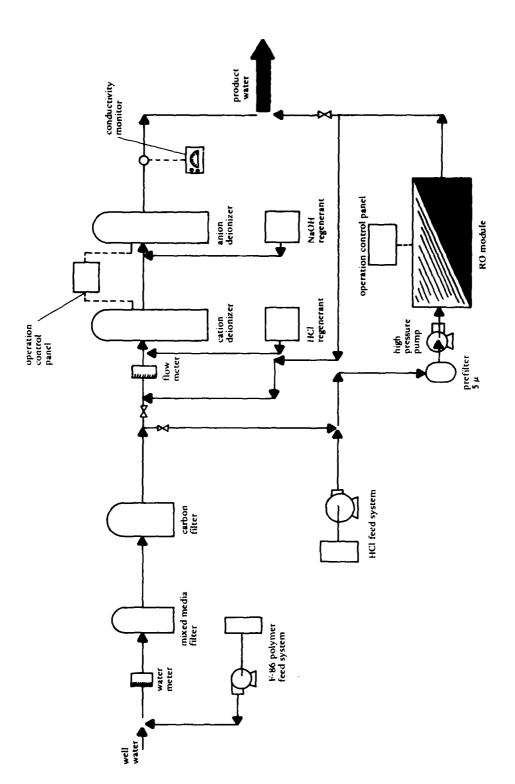


Figure 4. Water treatment system tested at Diego Garcia.

- 1. Monitoring the Well Water Quality
- 2. Identification of Pretreatment Requirements
- 3. Test of the Water Treatment System
- 4. Analysis of Water Samples for Total Trihalomethanes (TTHMs)
 Content
- 5. Sample Collection and Analysis
- 6. Milestones and Schedule

Monitoring the Well Water Quality

Individual wells were monitored daily to determine the amounts of TDS and chloride that were in the well water so that the impact on the water treatment system performance could be assessed. A sudden increase of TDS and/or chloride concentrations in well water means upconing and/or the intrusion of salt water. When chloride concentration was to exceed 250 mg/l, pumping would be terminated immediately in accordance with existing instructions.

The wells were monitored for additional water quality parameters at least once a week or as frequently as circumstances dictated. Grab water samples were considered satisfactory. The physical and chemical parameters tested and recorded are summarized and presented in Appendix B. A set of water quality field test kits was procured. They are listed below:

Test Kits and Parameters

| Test Kits* | <u>Parameters</u> |
|---|---|
| P-5085/2119 CWA/7611 TTM/7519 WAT-DC/4491 | pH Color Turbidity Alkalinity |
| CC-PS/4630 MPSC-DR/7932 EDO/7414 PHT-CM-DR/4824-DR | Sulfide Chloride Dissolved oxygen (DO) Total hardness, calcium hardness |

^{*}Procured from LaMotte Chemical Company.

Analytical procedures for using these kits were provided with each kit package. They are comparable to the HACH system and standard methods although their accuracies are somewhat lower. The kits were adequate for field use in terms of accuracy, time consumption, and manpower/skill requirements.

In addition, a TDS and dissolved solids (DS) test meter, made by Myron L. Meter Co., was used in the field. A Millipore Coli-Count Water Tester (MC 00 000 00) was used to test the coliform population in well and product water.

Identification of Pretreatment Requirements

The primary treatment objective is to reduce the TDS content from the current level of approximately 650 ppm to the EPA standard of below 500 ppm. In order to use the ion-exchange deionizer or the RO deminer-

alizer effectively, preliminary pretreatment is necessary.

Several different pretreatment tests were conducted: two filters, multi-media filter (first stage) and a carbon adsorption filter (second stage), were used in the water treatment system to provide the preliminary pretreatment of the raw water (or feed water) prior to its being processed through either an ion-exchange deionizer or RO demineralizer. The pretreatment of the feed water was designed to remove turbidity, suspended solids, color, taste, odor, excessive organics, and microorganisms in the water so that the product water would meet drinking water standards after deionization/demineralization. In the meantime, the ion-exchange resins and RO membrane would be protected through such pretreatment for a longer service time and useful life span.

Addition of acid to the RO system feed water was necessary in order to control scale formation on the RO membrane. As indicated previously, the pH of feed water must be maintained between 5.2 and 5.6, with 20°Be

Additional pretreatments were tested at the field site as alternatives and/or add-on processes for future applications. The pretreatments tested included addition of a polymer before filtration, an addition of NaOCl or $\rm H_2O_2$, or a combination of these two chemicals to oxidize organics and kill microorganisms.

Another treatment process, which could be used in place of multimedia filtration, is slow sand filtration. This process has long been recognized in water treatment as an effective means of removing suspended solids and microorganisms. The Naval Facilities Engineering Command recommended that NCEL test the process in the field. Accordingly, a small bench scale set-up, using coral sand, was tested to determine its applicability.

All of the pretreatment alternative tests were conducted at Diego

Garcia. The test results are presented in Appendix C.

Test of the Water Treatment System

The water treatment system consists of two mineral removal processes: a cation-anion ion-exchange deionizer and a RO demineralizer. Both components were tested for their effectiveness in removing TDS, chloride, hardness and other parameters. The reliability of operation and the requirements for maintenance were also evaluated. Test procedures for each of the water treatment systems are described as follows:

Ion-Exchange Deionizer. As indicated previously, a multi-media and a carbon adsorption filter was used for pretreatment of water to be fed into the ion-exchange deionizer. The deionizer has the capability of reducing mineral content (measured as TDS) in the feed water to 1 to 10 ppm. The ion-exchange capacity for the cation exchanger is about 230,000 grains expressed as CaCO_3 per regeneration cycle. A total of 30 regeneration cycles were run to determine the ion-exchanger capacity in the field situation. A mass balance was calculated to determine mineral removal effectiveness and efficiency of the ion exchanger. A Langelier Index (LI), for the ion-exchanger effluent, was calculated to assess the water's scaling or corrosion potential. Results of the deionizer operation are summarized and presented in Appendix D.

Reverse Osmosis Demineralizer. An RO demineralizer can remove not only the mineral content in the water but also other contaminants, such as organic chemicals and micro-organisms. During the test, the capability of the RO membrane to treat Diego Garcia water was thoroughly investigated. Fouling characteristics, the major concern for employing the RO process, was studied. To protect the membrane from fouling, pretreatment of the feed water was provided. A multi-media filter, a carbon adsorption filter, a polyelectrolyte feed an acid feed, and a 5 μ prefilter were the pretreatment components used in the RO water treatment system.

The polyelectrolyte F-86, a highly active cationic liquid polyelectrolyte, was used in conjunction with the multi-media filter to remove a majority of the colloidal particles (including microorganisms) in the water prior to using the RO system. These colloidal particles in the water were measured in terms of a fouling index (FI). Based on field experience and manufacturers' recommendation, the FI should be measured at the field operation site and be maintained below a value of 3 to avoid the frequent or premature cleaning of the RO membranes.

The pH control system, with a shutdown feature, was designed to continuously feed 20°Be HCl to the feed water of the RO system, and to shut off the acid feeder if the pH went below a designated preset range. By feeding acid to the feed water, the scale forming rate in the RO system was greatly reduced. The recommended pH setting ranges are between 5.2 and 5.6.

Methods and procedures for removing foulants from the RO membranes were provided in operational instructions. The modules were cleaned once during the entire test period to familiarize the operator with the cleaning procedure. A small amount of dirt appeared in the cleaning solution.

The RO system operated for more than 500 hours (cumulative time) during the test period. The recovery rate (ratio of total product water flow rate and feed water flow rate) was arbitrarily maintained at 70%. Definitions of operation parameters that should be noted on data sheets, such as recovery rate, salt passage rate, and pressure drop, were provided in the operating instructions manuals.

Data sheets for RO demineralizer's operations and water sample

analysis are presented in Appendix E.

Data were recorded daily and evaluated on a continuous basis. This evaluation and comparison helped in determining the consistency of RO system performance and frequency of cleaning. Evaluation techniques were provided in the operating instruction manuals. A graphic analysis that relates three parameters: salt passage rate, average pressure drop, and total product flow, was maintained for a better understanding of the RO system performance.

Analysis of Water Samples for Total Trihalomethanes (TTHMs) Content

On 29 November 1979, the EPA promulgated regulations limiting the permissible levels for TTHMs in drinking water (Ref 4). Recently, the American Water Works Association reached an agreement with the EPA on the TTHMs rule. A summary of the TTHMs regulations is described in Appendix F.

Removal and/or reduction of TTHMs in water can be achieved by:
(a) use of an oxidant disinfectant that does not generate trihalomethanes in water, or that generates less, (such oxidant disinfectants that can be used include: ozone, ozone-UV, chlorine dioxide, and chloramines); (b) treatment to reduce precursor concentrations prior to chlorination, such treatments include: off-line water storage, aeration, improved coagulation, ion exchange resins, granular-activated carbon (GAC), powdered-activated carbon (PAC), and ozone-enhanced biological activated carbon (BAC); or (c) treatment to remove TTHMs after formation, such treatments include: GAC, aeration, and macroreticular resins.

Impact of TTHM's Regulations to Diego Garcia. The regulations promulgated by EPA do not apply to Diego Garcia. BUMED does not provide any regulations or guidelines on TTHMs at this time (Ref 5). Nevertheless, the objective of the testing and monitoring of the TTHMs parameter during the test period was to ensure that TTHMs in the water is within the EPA limit and that the Fleet's readiness is adequately protected.

In the water treatment system, the second-stage carbon filter would remove the TTHMs and their precursors. The RO system provided has a molecular cuttoff weight of 200, therefore, any TTHMs with a molecular

weight of more than 200 could be removed by the RO process.

Collecting water samples for TTHMs analysis required a special sampling bottle and sampling procedure. TTHMs analytical methodology and procedure were provided by EPA (Ref 4). A gas chromatopgraphy/mass spectroscopy instrument was used for quantitative and qualitative analysis of TTHMs in a water sample. The Chemistry Laboratory at the Public Works Center (PWC), Guam, has the capability and facility for TTHMs water sample analysis. Four series of water samples collected at Diego Garcia were sent to PWC, Guam, for TTHMs analysis. The results are presented in Appendix G.

Sample Collection and Analysis

Water samples were collected and analyzed as follows:

- 1. During system startup daily.
- 2. During test period twice weekly.
- 3. Occasionally, a coliform count for raw and product water was performed.

- 4. Every two weeks water samples were collected at Diego Garcia along with one of the biweekly sample collections, sent to PWC, Subic Bay, for analysis of all physical, chemical, and biological parameters. In addition, a chemical oxygen demand (COD) analysis was planned, but due to transportation problems, only one set of water samples was sent to PWC, Subic Bay. The test results have not been received from PWC, Subic Bay.
- 5. Four series of water samples were collected for TTHMs analysis. (The water samples were sent to PWC, Guam for analyzing.)

Milestones and Schedule

The milestones and test and evaluation schedule of the water treatment system at Diego Garcia for 1981 are shown below:

| Milestones | March | April | May | June | July | August |
|--|-------|-------|-----|------|-----------|--------|
| System Installation Well Water Quality | 0-0 | 0 | | | | |
| Water System Test | Ω | | | | 0 | |
| Data Collection | D | | | | 0 | |
| Data Analysis | C) | | | | | |
| System Disposition | | | | | - | 0 |

DISCUSSION OF TEST RESULTS

The test results on well water quality, pretreatment requirements, water treatment system operation, and TTHMs content analysis are presented in Appendixes B, C, D, E, and G. The results and findings are discussed below:

Well Water Quality

Blended well water from about 14 wells was used in the test and evaluation of the water treatment system. Because a parameter affecting the quality of the blended water was an average of all the individual wells, a slight increase could be due to a large increase of that parameter in a single well. Therefore, daily well water samples from each well were checked for TDS and chloride concentration. Any well with chloride exceeding 250 mg/l would be shut down or operated at a reduced pumping rate.

The values of other parameters previously discussed plus alkalinity, dissolved oxygen, sulfide, and pH were determined on a weekly basis or more frequently if needed. These additional parameters were measured for assessment of well water quality. The well water quality did not change much. However, a few wells, such as C-2 and C-3, in which C-2 had been secured for some time due to a high conductivity (or TDS) reading, did vary significantly in quality (see Appendix B).

In general, the pH values of the well water ranged from $6.8\ \text{tc}\ 7.7$, the raw water had color and turbidity readings exceeding EPA standards and had a high alkalinity content (300 to 850 ppm of CaCO_3) due to high bicarbonate and associated hardness content. The chloride content ranged from 24 to 456 ppm (average composite was 100 ppm), while TDS concentrations ranged from 300 to 1400 ppm (average composite was 600 ppm). Alkalinity values were consistently higher than TDS.

As reported in the Background section, the composite well water quality generally met EPA's standards except for TDS, turbidity, and color. Among these parameters, the turbidity measurement might have been affected by the color content in the water. It is anticipated that ultraviolet radiation units now being installed, and existing aeration and chlorination are adequate to treat the composited well water for

potable use.

Pretreatment Requirements

There were three components in the water treatment system that provide pretreatment of the well water to prevent fouling of the desalination unit, either the ion-exchange deionizer or the RO demineralizer. The components are a polyelectrolyte, a multi-media filter (first stage), and a carbon filter (second stage). The pretreatment not only prepared water suitable for desalination, but also provided a treatment for removing turbidity, microorganisms, color, taste, odor, and organics (including TTHMs formation precursors). These pretreatments produced water meeting a portion of EPA standards and made it more palatable. The efficiencies of these pretreatments are presented in the following sections together with the water treatment system.

The water fed into the RO system required two additional treatments to the pretreatments just mentioned. These treatments were a pH adjustment and a 5 μ cartridge filter. The pH adjustment, in terms of hydrochloric acid requirement, is discussed below. Titration data are shown

in Appendix C.

<u>pH Adjustment.</u> Titration experiments were conducted at Diego Garcia to determine what dosages of hydrochloric acid were needed to adjust pH in the feed water. The strength of the hydrochloric acid used was one-tenth of 20° Be (31.4%) HCl stock solution. It was found that in the laboratory titration experiment, an average of 0.045% (volume by volume, (v/v)) of hydrochloric acid was required to adjust the feed water's pH to 5.6. This correlated well with the RO operational data, in which 0.057% (v/v) was the average dosage required. Acids available for use in the RO process are described in Appendix H.

Slow Sand Filtration. A slow sand filter, fabricated at Diego Garcia, was tested and evaluated to determine its capability to remove water contaminants. The filter had the following characteristics:

Filter size: 5 1/8 in. diam by 6 ft long

Sand depth: 42 in. (coral sand media)

Gravel depth: 6 in. (coral gravel)

Sand media: Effective size (0.5 mm), uniformity

coefficient (1.5)

Filtration rate 0.07 gpm/ft²

The results of 16 days of continuous operation indicated that the water's color was consistently reduced from 20-30 to 5 except the last 2 days (reduced to 10 only) and the turbidity from 3 to 0. The water quality would meet EPA standards for these two parameters. However, the flow rate of the filter dropped from 35 ml/min (0.07 gpm/ft² equivalent surface loading rate) controlled effluent flow rate to 15 ml/min (0.03 gpm/ft² equivalent surface loading rate) effluent flow rate uncontrolled, i.e., the valve was wide open. This reduced flow rate is an indication of clogging. Further investigation indicated that the sand filter had a limited capacity for removing the bacteria cells, even with the addition of disinfectants, i.e., H_2O_2 and/or Ca(ClO)₂.

Fouling Index (FI)

When water contains particulates smaller in size than can be removed by the available filter media, it is necessary to increase this particle size in some way. Coagulation is the most common method, and aluminum sulfate has been used as a coagulant for many years. The process requires large settling tanks and long retention times. Recently polyelectrolyte compounds have been introduced which accomplish the coagulation more rapidly. As mentioned earlier, long membrane life requires a water with minimum amounts of particulates. An indicator of filtration efficiency, FI, is used where high quality water is required. FI is defined as follows:

FI =
$$\frac{\text{Final Time } (T_f) - \text{Initial Time } (T_i)}{T_f} \times 10$$

where T_i = time required to collect 500 ml of a water sample filtered through a Millipore filter at the beginning of test (sec)

 T_f = time required to collect 500 ml of a water sample filtered through the same Millipore filter after the filter is continuously used for 10 min (sec)

Millipore filter used: 5HA 0.45 μ Filtration pressure used: 30 psi

The manufacturer of the hollow-fiber membrane recommended an FI of 3.0 or less for good membrane performance and also recommended a cationic liquid polyelectrolyte (F-86) being used at Diego Garcia for controlling FI.

The amount of F-86 to use was arrived at by trial and error. The correct amount of F-86 to use was obtained by gradually increasing the F-86 feed until the FI reduction leveled off and then began to increase. At Diego Garcia the FI of the raw feed water was in excess of 8. The F-86 dosage was started at about 15 ppm (v/v) and increased incrementally with only a slight reduction of the FI; at 20 ppm the FI was 7.5.

Increasing the retention time between the point of F-86 injection and the filter was tried as a possible means of improving the FI. This was accomplished by adding a 100-gal. tank in the line, and also by reducing the flow to one-half and one-fourth the normal rate. No appreciable reduction in FI was obtained.

The high concentration of F-86 did clog the mixed media filter to the point that it could not be backwashed effectively. The entire anthracite portion of the filter was a solid mass with a few passages through it where the flow channeled. Physical breakup of the mass was not enough to assist in the backwash. Laboratory field tests indicated that NaOH would remove the clog, so 50% NaOH was added to the filter with enough water to make a 12.5% solution. A series of soakings and backwashes finally returned the filter to its original state. Fifteen days later the same condition was found and again the filter was cleaned with NaOH. At this time it was decided to discontinue using F-86. Some other electrolyte might work better at Diego Garcia.

Culligan commented on the FI results after the field test was completed. They recommended that the FI also be measured at the prefilter (5 μ cartridge filter) effluent. This was the final treatment that the water was subjected to before entering the RO system. This means that the prefilter would be used as a "sacrificer" to ensure that the membrane would be protected. This appeared to be the case at Diego Garcia because using F-86 and the multi-media filter treatment failed to reduce the FI value to 3.0 and below. However, the membrane appeared to be protected, because the membrane performed well according to the data (recovery rate and salt passage percentages) shown in Appendix E. Replacement of the prefilter toward the end of the field tests supports the assumption that the prefilter was sacrificed.

Even if a satisfactory solution can be found to control the FI, it is considered advisable to use a spiral wound membrane which is less liable to fouling, is easier to clean than the hollow-fiber filter, and will not increase the cost. One manufacturer of the spiral wound membrane claimed that fouling is not a serious problem. They have projected a 3-year life for their membrane, based on operating data and water analyses.

Chemical Additions

Two experiments were conducted at Diego Garcia to determine the feasibility of adding $Ca(0Cl)_2$ and H_2O_2 for reducing color, odor, H_2S_3 , and bacteria count and to increase dissolved oxygen. The results shown in Appendix C were not as good as had been anticipated. Logistic support for either treatment would not be justified for the limited benefit obtained.

Water Treatment System

As stated previously, the water treatment system consists of a three-stage process: multi-media filter, carbon filter, and desalination. The first and second stages of the treatment did remove or reduce the color, turbidity, taste, odor, and most organic chemicals in the well water so that a portion of the EPA standards were met. The third stage, desalination, either by ion-exchange or reverse osmosis, is discussed as follows:

Ion-Exchange Deionizer. As shown in Appendix D, the two-bed cation and anion exchanger effectively removed TDS down to 4 to 42 ppm range (measured by conductivity), total hardness to 0 to 16 ppm, calcium to 0 to 15 ppm, chloride to 7 to 76 ppm, and alkalinity to 100 to 300 ppm. The product water quality met the EPA requirement of 500 ppm TDS and 250 ppm chloride. The effluent water quality was measured continuously by a conductivity cell (measurement of TDS) connected to a meter with the capability of stopping the operation when a preset conductivity level was reached. Since the capacity of the resin beds is fixed with respect to the concentration in ppm of ion removal, the quantity of product made and the time of each cycle is indirectly proportional to the TDS of the feed water. For example, during the first hour or two of operation on a newly regenerated ion exchange resin on 500 ppm feed, the product was as low as 1 ppm TDS. For the major portion of the cycle, the TDS level was about 12 to 15 ppm. When the resin was exhausted, the TDS climbed very rapidly to the preset conductivity level, shutting down the unit. When the deionizer was operated on effluent water from the RO unit (40 ppm TDS), the product water was about 10 to 12 ppm during the major portion of the run. Cycle times were much longer, although the actual figures were not determined.

During the test period, 1,040 gallons of 20°Be HCl (31.4%) and 285 gallons of NaOH (50%) were used to regenerate the resins while producing 492,290 gallons of high quality water. This would require 2.11 gallons of HCl and 0.579 gallon of NaOH per 1,000 gallons of product water. It is estimated that a 450,000-gallon water treatment plant would require 950 gallons of HCl and 260 gallons of NaOH per day for resin regeneration. This would create a considerable logistic

support problem.

The amount of chemicals would be drastically reduced, of course, if the deionized water was blended with the pretreated high TDS water to obtain a product having less than 500 ppm TDS. For example, if 1,000 gallons of 20 ppm TDS water were added to 4,800 gallons of 600 ppm TDS water this would provide 5,800 gallons of water at 500 ppm TDS. This would be equivalent to 0.364 gallons of HCl and 0.1 gallon of NaOH/1,000 gal product water at 500 ppm TDS. The 450,000-gallon plant would require 163 gallons of HCl and 45 gallons of NaOH; therefore, the minimum cost for chemicals would be approximately \$1.00/1,000-gal product water (FOB Diego Garcia based on a price factor of 3). When ${\rm H_2SO_4}$ is substituted for HCl the cost is reduced to \$0.60/1,000-gal product water. For a 250 ppm TDS water product the above costs would be \$3.50 and \$2.10, respectively.

Reverse Osmosis Demineralizer. The RO demineralizer and its first and second stage filters were the major treatment processes tested and evaluated at Diego Garcia. The performance of the RO system was considered highly satisfactory. The product water quality far exceeded EPA standards.

In reviewing the data obtained during the RO operation (Appendix E), the RO membrane was capable of removing about 90% of TDS in the feed water, while maintaining a 70% recovery rate. Although the FI remained high because of the inability to control it with the polyelectrolyte, F-86, the RO system's performance did not appear to be affected. The prefilter apparently served as a "sacrificer" for assuring RO feed water quality in terms of FI value. The cost of a prefilter cartridge is \$4.75 each. Using FOB Diego Garcia based on a price factor of 3, the prefilter will cost \$71.25 (or 5 at \$14.25 each), which is considered relatively inexpensive. Furthermore, in reviewing the percentage of salt passage (ratio between effluent TDS ppm to the total TDS ppm in feed water), the membrane performed well during the entire test period.

Towards the end of the field test, brackish water was simulated by adding seawater to bring the TDS to 1,000, 1,500 and 2,000 ppm (See Appendix E). A 70% recovery rate and a 6 to 8% salt passage rate (or 94 to 92% salt rejection rate) were maintained. Though the TDS concentration in the product water had increased, the EPA standards were still met. This finding indicates the RO system can operate on a high concentration of TDS without altering the operational procedure or sacrificing the production rate.

The acid (20°Be 31.4% HCl) consumption rate was estimated at Diego Garcia at 0.057% (v/v) for pH adjustment. This is equivalent to 0.81 gallon HCl (20°Be 31.4%)/1,000 gallons of product water at 70% recovery rate. For a 450,000-gallon water treatment plant, 366 gallons of HCl will be required. The logistic support for handling this much acid could be significant. As calculated in Appendix H, when 94% concentration H_2SO_4 is used, the volume of the acid can be significantly reduced to 103^2 gallons.

Chemical consumption rates between the ion-exchange deionizer and the RO demineralizer are compared as follows:

| Process | Product | Water | Raw Wa (mix wi | | | . Rates | |
|---------------------------|-------------------|--------------|-------------------|--------------|--------------|---|-------|
| Frocess | Volume (K gal) | TDS (ppm) | Volume (K gal) | TDS (ppm) | HC1 (gal) | H ₂ SO ₄ (gal) | (gal) |
| Ion-Exchange Deionizer | 77.59 | 20 | 372.41 | 600 | 164 | 46 | 45 |
| RO Demineralizer | 88.33 | 60 | 366.67 | 600 | 67 | 19 | None |

The 450,000 gallons of blended product water will contain 500 ppm TDS from original raw water containing 600 ppm TDS.

The concentration of chloride (when HCl is used) or sulfate (when H2SO₄ is used) in product water must be monitored to assure that the EPA standards are met. Power requirement was estimated at 12.0 KWH per 1,000 gallons of product water.

A preliminary design of a 300,000-gpd (product water) RO system is illustrated in Appendix E. Better salt balance data (cation and anion analysis) shall be provided for a closer preliminary design of a full-

scale RO system.

Total Trihalomethanes (TTHMs) Concentration in the Water

Four series of water samples were collected for TTHMs concentration analysis by PWC, Guam. The results (Appendix G), indicated that the TTHMs concentration in the water at Diego Garcia appeared insignificant. In the second and fourth series of samples, though, there were two samples, no. 2 multi-media effluent and no. 3 carbon filter effluent which exceeded the EPA primary standard of 100 ppb, yet with a follow-on treatment with either the ion-deionizer or the RO system the TTHMs concentration was reduced to below 10 ppb. The carbon filter supposedly would adsorb TTHMs and its precursors. That the carbon filter effluent contained higher TTHMs concentration than raw well water and multi-media filter effluent can only be explained by: (a) the activated carbon was exhausted in adsorbing TTHMs, (b) there was suddenly a release of TTHMs previously adsorbed on carbon, and/or (c) a bio-discharge occurred due to sloughing-off or biomass disintegration, and (d) laboratory analytical errors.

Low TTHMs concentration in the ion-exchange effluent appeared strange, unless the TTHMs precursors were in radical forms that could be temporarily ion-exchanged. As for the RO membrane, should there be a molecular weight cut-off at 200 (as claimed by the membrane manufacturers), then two of the TTHMs species, CHClBr₂, and CHBr₃ should have been removed. In natural well water, however, the formation of brominated compounds is less probable than chlorinated compounds. Therefore, it appears that there is a need to review the accuracy of the analytical procedure.

The first series of samples taken by a technician of PWC, Guam, appeared to have a high concentration of TTHMs, but no satisfactory

explanation could be provided.

WATER QUALITY REQUIREMENT

The desirable characteristics of water vary with its intended use. Water for drinking and food preparation must be free from organisms capable of causing disease, and from minerals and organic substances producing adverse physiological effects. To encourage man to drink this health-promoting liquid, the water must also be aesthetically acceptable. For example, the water should be free from turbidity, color, taste, and odor. Drinking water should also have a reasonable temperature. The term "potable water," means that it can be consumed in any desired amount without concern for adverse effects on human health.

The water produced by the experimental water treatment system at Diego Garcia betters EPA's standards in terms of turbidity, color, taste, and odor, chloride, TDS, microorganisms, and organic substances. However, the water is not suitable for long term consumption primarily due to the low mineral content (TDS 50 ppm) and low pH value (about 5.5), i.e., the water is "too pure and too acidic." This type of water can cause severe corrosion problems in metal pipelines and is generally used in medical and general chemistry and certain industrial operations, e.g., electronics, photo labs, boiler/cooling towers. Blending the demineralized water with some of the raw well water processed from the two filters will provide a suitable potable water.

Langelier's Saturation Index

Langelier's Saturation Index (LI) is the most widely used method to measure the water quality for scaling or corrosion potential. Appendix I provides the LI calculation formula.

Using water quality data obtained at Diego Garcia, one can easily calculate the LI at which scale or corrosion can be predicted.

For example:

Date: May 11, 1981. Ion-Exchange Deionizer Operation, Temp = 28°C

| | TDS | Total <u>Hardness</u> | Ca <u>Hardness</u> | <u> </u> | рН |
|---------------|-----|--------------------------|-----------------------|----------|-----|
| Well Water | 460 | 180 | 104 | 550 | 7.7 |
| Product Water | 10 | 16 | 4 | 200 | 5.5 |

LI Well Water = (+) 0.65 (scale formation) LI Product Water = (-) 3.28 (corrosion)

Date: May 9, 1981 Reverse Osmosis Operation, Temp = 28°C

| | TDS | Total <u>Hardness</u> | Ca <u>Hardness</u> | Alk | рН |
|---------------|-----|--------------------------|-----------------------|-----|-----|
| Well Water | 440 | 184 | 112 | 650 | 7.6 |
| Product Water | 44 | 4 | 0 | 200 | 5.5 |

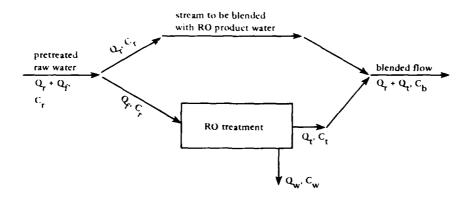
LI Well Water = (+) 0.6 (scale formation) LI Product Water = (-) 2.22 (corrosion)

By using the same equation, one can also calculate an adequate pH level to be maintained (or adjusted) in the RO feed water so that calcium or magnesium carbonate (scale) deposits will not form on the RO membrane. For this purpose, LI is recommended by the membrane manufacturer to be about -0.2, which is slightly corrosive. A calculation is illustrated as follows (using May 9, 1981 well water quality data):

This theoretical pH value appeared to be quite different from the manufacturer's recommended actual field operation value of 5.6. This recommendation was made to compensate for any drastic changes in well water quality (e.g., a sudden increase in hardness). Additional testing will be required to determine what will be the most appropriate pH level. The acid required for pH adjustment can be reduced by two-thirds when pH 6.7 RO feed water is allowed, rather than 5.6 (refer to pH adjustment data, Appendix C (1)).

To use the RO system product water as drinking water, the pH must be readjusted to at least 6.5 and mineral content be increased to 250 ppm TDS (optimum). Sodium bicarbonate is the most commonly used chemical for such purposes. Other methods that can be used to solve the problem without any cost are (a) split treatment and blending, and (b) the use of a coarser and newer membrane that will require lower power/pressure and produce water with higher TDS.

Calculation of Split Treatment Requirement



Where:

 Q_r = raw water flow rate, (to be blended with treated water), L^3/T

 Q_f = feed water flow rate, (to be treated with RO), L³/T

 Q_t = treated water flow rate, (considering certain waste of feed water), L^3/T

 $Q_{...}$ = waste water stream from treatment process, L^3/T

 $C_m = total$ dissolved solids (TDS) concentration in raw water, M/L^3

 $C_b = total$ dissolved solids (TDS) concentration in blended water, M/L^3

 $C_t = total dissolved solids (TDS) concentration in treated water, <math>M/L^3$

 $C_{W} = total dissolved solids (TDS) concentration in waste stream, M/L³$

Define:

$$R = Q_t/Q_f \text{ (RO treatment recovery rate) or } Q_t = RQ_f$$

$$r = Q_f/Q_f+Q_r \text{ or } Q_r = (1-r)/r Q_f$$

$$p = 1-C_t/C_r \text{ (TDS removal efficiency), or } C_t = C_r(1-p)$$

Mass balance:

$$C_{r}Q_{r} = C_{b}Q_{b} - C_{t}Q_{t}$$

$$C_{r}Q_{r} = C_{b}(Q_{r}+Q_{t}) - C_{f}(1-P)(Q_{f}R)$$

$$Q_{r} = Q_{f}R [C_{b} - C_{f} (1-P)]/C_{f} - C_{b})$$
(1)

Blended Water Treatment

Equation (1) shows the flow rate of pretreated raw water to be blended with RO treated water.

For example, a demand of 450,000 gallons of 500 ppm TDS water for Diego Garcia would require 485,000 gallons of 600 ppm TDS well water fed into the pretreatment system. The pretreated water is then split into two streams: 118,000 gallons is processed through the RO system to produce 83,000 gallons of 60 ppm TDS water, assuming a 70% recovery rate. The other 367,000 gallons of water, with no further treatment, is blended with the 83,000 gallons of RO product water. This equation can be further developed to provide the relationship between TDS concentrations in raw water and blended water. For example:

$$(1-r)/r Q_{f} = Q_{f}R [C_{r}(1-p) - C_{b}]/(C_{b}-C_{r})$$

$$(1-r) (C_{b}-C_{r}) = rR [C_{r}(1-p) - C_{b}]$$

$$(1-r)C_{b} - (1-r)C_{r} = rR (1-p) C_{r} - rRC_{b}$$

$$C_{b} = [(1-r) + rR(1-p)]C_{r}/(1-r) + rR$$

$$(2)$$

Concentration of TDS in the waste stream can also be estimated as follows:

$$Q_{f}C_{r} = Q_{t}C_{t} + Q_{w}C_{w}$$

$$Q_{t}/R C_{r} = Q_{t}C_{r} (1-p) + (Q_{t}/R - Q_{t}) C_{w}$$

$$C_{r} = C_{r}R (1-p) + (1-R) C_{w}$$

$$C_{w} = [R (1-p) - 1]C_{r}/(1-R)$$
(3)

COST COMPARISON FOR DESALINATION PROCESSES

Desalination processes include distillation, reverse osmosis, electrodialysis, ion-exchange, and freezing. Distillation has been used for seawater conversion for many years, but is not economical to treat brackish water. Recent high energy and skilled personnel costs have made distillation also less economical for seawater. RO has been used successfully on brackish water for about 10 years and within the past 5 years the technology has been used to economically treat seawater for potable use. Electrodialysis (ED) and ion-exchange (IX) methods are generally used for treating brackish water only. The freezing method is still in the development stage; its applicability in the field still needs to be proved.

Economical comparison of the desalination processes is very difficult since they are generally site specific. Pretreatment requirements of feed water for these processes has a significant impact on the costs.

Based on studies published in AWWA journals (Ref 5 and 6), the RO process has become the least-cost option for demineralization of the bulk of brackish water supplies in the western United States.

The water supply capacity of the 15 communities studied (see Tables 1 through 4), ranged from 0.15 to 6.10 MGD. The TDS concentration in the feed water was between 941 to 3236 ppm. The demineralization cost in 1977 dollars ranged from \$0.37 per 1,000 gallons of water to \$1.56 per 1,000 gallons of water. The study revealed that in seven of the municipalities, RO is projected as the least-cost demineralizing system; in six, RO is combined with ion-exchange or zeolite deionizer (Z); and in the remaining two, RO and ED are considered an even choice. Recent developments in the RO membrane technology, as well as the mass production of the membranes, has not only reduced the RO system's capital investment and O&M cost, but has also made it more reliable and favorable for future applications. For instance, the RO manufacturers have lowered the feed water pressure requirements from 800 to 600 to 400 psi while maintaining membrane permeation rates.

Optimization of RO system designs to employ looser membranes - having somewhat lower salt-rejection characteristics, yet yielding potable water from high salinity feeds at lower pressures - is one objective of current RO manufacturers. Such opportunities do not appear to exist for lowering the energy requirements of ED systems.

Table 1
Projected Demineralization Costs for 15 Candidate Communities, 1970-1973

| | | | | | === | | | | | | |
|--------------------------|----------------|------|---------------|------|--------------------------|--------|----------------------------|---------------|-------|---------------|--|
| | | w | water | | icipal ater fity ~ | | Demineral- ization Cost | | | | |
| | Popu- | mg/l | | mg/f | | Demin- | 70-73* | | 1977* | | |
| Munici- pality | lation 1970 | TUS | Hard- ness | TUS | Hard- ness | | €/m² | c/1000 gal | €/m* | e/*100 gal | |
| Lakota, N. D. | 964 | 1754 | 816 | 195 | 100 | IX-RO | 41.5 | 157 | 45.2 | 171 | |
| Eureka, S. D. | 1 547 | 2163 | 1248 | 430 | 5 | Z-RO | 20 | 75 | 30.1 | 114 | |
| Sibley, Iowa | 2 747 | 2720 | 1547 | 481 | 100 | IX-ED | 38.3 | 145 | 50.4 | 191 | |
| Freer, Tex. | 3 000 | 1242 | 77 | 500 | 49 | ED I | 15 | 56 | 27 7 | 105 | |
| Towanda, | 1 242 | 2417 | 1033 | 246 | 125 | RO | 36.2 | 137 | 55 2 | 200 | |
| Kan. | | } | 1 | ļ | } | 1 | | | | l l | |
| Roundup. | 2 010 | 1643 | 725 | 385 | 72 | RO | 20 | 77 | 34 3 | 130 | |
| Malta, Mont. | 2 195 | 969 | 332 | 493 | 29 | l IX | 11 | 42 | 19 | 71 | |
| Las Animas, Colo. | 3 100 | 3236 | 1512 | 981 | 194 | RO | 15 | 57 | 25 | 96 | |
| Fl. Lupton, Colo. | 2 500 | 1423 | 588 | 130 | 200 | RO | 17 | 42 | 18 | 89 | |
| Ft. Stockton. Tex. | 7 500 | 1796 | 798 | 485 | 150 | IX-ED | 10 | 40 | 20 | 75 | |
| Kihei, Haw. | 13 500 | 1100 | 207 | 500 | 90 | ED | 15 | 57 | 20 | 75 | |
| Cesa Grande, Ariz. | 10 500 | 945 | 380 | 500 | 205 | ED | 8.2 | 31 | 13 | 50 | |
| Midland. Tex.t | 63 000 | 1135 | 503 | 400 | 100 | IX-RO | 6.9 | 26 | 11 | 42 | |
| Ark. City, Kan. | 14 000 | 1658 | 565 | 500 | 65 | ED | 9.2 | 35 | 14 | 53 | |
| Artesia. N. M. | 10 315 | 941 | 853 | 490 | 100 | ix | 9.0 | 34 | 15 | 57 | |

*1970-1973 costs equated to 1977 dollar values iFor supplement to surface water supply

Table 2

Demineralization Plant Flows, 1970-1973

| -: | Fend | | Demineral- ization | | Brine | | Blend | | Supply | |
|----------------|------------|------|-----------------------|------|--------|------|---------|------|--------|------|
| Stunicipality | ın'/day | myd | m'/day | mgd | m /day | mgd | nt'/day | med | m:/day | mgd |
| Likota, N. D. | 790 | 0.21 | 570 | 0 15 | 200 | 0.06 | 0 | 0 | 570 | 0 15 |
| irka, S D. | 1 300 | 0.35 | 3 100 | 0.30 | 200 | 0.05 | a | 0 | 1 100 | 0.30 |
| Shirv, lowa | 1 900 | 0.50 | 1 400 | 0.36 | 5:10 | 0 14 | 0 | 0 | 1400 | 0.36 |
| Freer, Tex. | 1 800 | 0.47 | 1.500 | 0.40 | 300 | 0 07 | 0 | ก | 1 500 | 0.40 |
| Juwanda. | 2 700 | 0.71 | 1 900 | 0 49 | 790 | 0 21 | 40 | 0.01 | 1900 | 0 50 |
| Kan. | | i | 1 | } | | 1 | | | l i | |
| Roundup. | 2,790 | 0.71 | 2 400 | 0.64 | 300 | 0.07 | 0 | 0 | 2 400 | 0.64 |
| Mont. | [| • | İ | i | 1 1 | ľ | | | | |
| Walta, Mont. | 3 500 | 0.92 | 3 000 | 0.79 | 490 | 0 13 | 0 | 0 | 3 000 | 0.79 |
| s Animas, ويرا | 8 330 | 2.20 | 5 300 | 1.40 | 3 000 | 0.80 | 0 | 0 | 5 300 | 1 40 |
| Colo. | l i | | Ì | 1 | 1 | | 1 | | i : | |
| Ft. Lupton. | 7 190 | 1.90 | 6 060 | 1.60 | 1 100 | 0.30 | 0 | 0 | 6 060 | 1.80 |
| Colo. | } . | ŀ., | | 1 |) : | | | 1 | | i |
| Ft Stockton, | 10 600 | 2.80 | 9 460 | 2.50 | 1 100 | 0.30 | 0 | 0 | 9 460 | 2.50 |
| Tex. | | · · | i | (| [1 | | 1 | | i i | l |
| Kihei, Haw. | 13 000 | 3.45 | 11 400 | 300 | 1 700 | 0.45 | 0 | 0 | 31 400 | 3.00 |
| Casa Grande, | 15 700 | 4.15 | 11 000 | 2.90 | 950 | 0.25 | 3 790 | 1.00 | 14 800 | 3.90 |
| Ariz. | 1 | | 1 | 1 | 1 | | 1 | | | ĺ |
| Midland, Tex. | 21 200 | 5.60 | 18 900 | 5 00 | 2 300 | 0.60 | 0 | 0 | 18 900 | 5 00 |
| Arkansas City. | 21 800 | 5.75 | 18 900 | 500 | 2 800 | 0.75 | 0 | 0 | 18 900 | 5.00 |
| Kan. |) | l | | 1 | 1 | | 1 | | [| 1 |
| Artesia, N. M. | 25 000 | 6.60 | 20 100 | 5.30 | 1900 | 0.50 | 3 000 | 0.6 | 23 100 | 6.10 |

*For supplement to surface water supply

Table 3
Projected Demineralization Costs for Fifteen Candidate
Communities, 1977

| | Municipal Water Quality- | | | Demineralization Cost | | | | |
|---------------------|--------------------------------|----------|----------------------------------|--------------------------|---------------|------|---------------|--|
| | | mg/l | D | 1970 | -1973* | 1977 | | |
| Municipality | TDS | Hardness | Demineral- ization Process | e/m' | e/1000 gal | s/m | ¢/1000 gal | |
| Lakota, N.D. | 157 | 61 | IX-RO | 45.2 | 171 | 412 | 156 | |
| Eureka, S.D. | 456 | 5 | Z-RO | 30 1 | 114 | 21 | 80 | |
| Sibley, lowa | 263 | 148 | IX-RO | 50.4 | 19: | 22 | 82 | |
| Freer, Tex | 341 | 21 | RO | 27 7 | 105 | 21 | 79 | |
| Towanda, Kan. | 500 | 42 | IX-RO | 55.2 | 209 | 18 | 67 | |
| Roundup, Mont. | 267 | 76 | IX-RO | 34.3 | 130 | 20 | 75 | |
| Malta, Mont | 261 | 90 | RO | 19 | 71 | 18 | 69 | |
| Las Animas, Colo. | 363 | 145 | IX-RO | 25 | 96 | 17 | 63 | |
| Ft. Lupton, Colo. | 258 | 107 | RO | 18 | 69 | 16 | 60 | |
| Ft. Stockton, Tex. | 270 | 120 | RO | 20 | 75 | 17 | 56 | |
| Kihei, Haw. | 300 | 94 | RO | 20 | 75 | l m | 41 | |
| | 1 | { | ED or | l | (| { | (· · · | |
| Casa Grande, Ariz. | 220 | 89 | RO | 13 | \$0 | 13 | 49 | |
| | 1 | 1 | 1X-ED or | 1 | 1 | 1 | 1 | |
| Midland, Tex.† | 370 | 100 | IX-RO | 111 | 42 | 9.8 | 37 | |
| Arkansas City, Kan- | 323 | 91 | RO | 14 | 53 | 14 | 52 | |
| Artesia, N.M. | 160 | 111 | RO | 15 | 57 | 13 | 48 | |

^{*1970-1973} costs equated to 1977 dollar values †For supplement to surface water supply at Midland

Table 4
Demineralization Plant Flows, 1977

| | Feed | | Demineral- ization | | Brine | | Blend | | Supply | |
|------------------------|----------|------|-----------------------|------|--------|-------|--------|------|--------|-------|
| Municipality | m /day | nigd | m'/day | mgd | m·/day | nigd | m /day | mgd | m vday | mgd |
| Lakota, N.D. | 490 | 0.13 | 380 | 0.10 | 110 | 0.03 | 190 | 0.05 | 570 | 0.15 |
| Eureka, S.D. | 1 300 | 0.34 | 1 100 | 0 30 | 150 | 0.04 | 0 | 0 | 1 100 | 0.30 |
| Sibley, lowa | 1700 | 0.45 | 1 400 | 0.36 | 300 | 0.09 | i o i | 0 | 1 400 | 0.36 |
| Freer, Tex. | 1700 | 0.44 | 1 500 | 0.40 | 150 | 004 | 0 | 0 | 1 500 | 0 40 |
| Towande, Kan | 2 200 | 0.58 | 1900 | 0.50 | 300 | 0 04 | 0 | 0 | 1 900 | 0 50 |
| Roundup, Munt. | 2.700 | 0.71 | 2 400 | 0.64 | 260 | 0.07 | ٥ | 0 | 2 400 | 0 64 |
| Malta, Mont | 3 300 | 0.88 | 3 000 | 0.79 | 300 | 0.09 | 0 | 0 | 3 000 | 0 79 |
| Las Animas. Colo. | 6 700 | 1.77 | 5 300 | 1.40 | 1400 | 0 37 | 0 | 0 | 5 300 | 1.40 |
| Ft Lupton. Colo | 7 040 | 1 86 | 6 OHO | 1 60 | 986 | 0.26 | 0 | 0 | 6 060 | 1 60 |
| Ft. Stockton, Tex | 11 400 | 3 00 | 9 460 | ₹ 50 | 1900 | 0 50 | υ | 0 | 9 460 | 2.50 |
| Kiher, Haw | 12 300 | 3 24 | # 330 ¹ | 2.20 | 910 | 0 24 | 34480 | 080 | 31 400 | 3 110 |
| Casa Grande, Aria | 16 (410) | 4 38 | 14 800 | 3 90 | 18110 | 0.48 | 0 | a | 14 600 | .1 90 |
| Midland, Tex. | 20 600 | 5 44 | 18 900 | Som | 1700 | 10 44 | 0 | | 18 900 | 5 00 |
| Arkansas City. Kari | 21 8681 | 5.75 | OCF: NS | 5 00 | 2641 | บ 75 | 0 | ď | 18 900 | 5 00 |
| Arresta, N.M. | 27 200 | 7 18 | 23 toU | 6 10 | 4090 | t con | 0 | o . | 23 100 | e 10 |

For applement to surface water supply

The brackish water RO membrane can take feed water containing TDS up to 15,000 ppm. The saline water RO membrane, however, will treat from 15,000 up to 50,000 ppm TDS. Since the pressure required for an RO system is directly proportional to the TDS content in the water (about 10 ppm TDS/psi max), it will be cost effective to use brackish water rather than seawater. Generally, the cost for using the RO system for treating brackish water is about one-third the cost of treating seawater.

FINDINGS AND CONCLUSIONS

- 1. The water treatment system tested at Diego Garcia accomplished the objective of reducing TDS turbidity, hardness, chloride, color, taste, and odor.
- 2. Several tests made on synthetic brackish water indicated that similar reductions could be attained to meet the standards with little change in the production rate or chemical demand.
- 3. The applicability of a slow sand filter at Diego Garcia is limited in terms of pretreatment for minimizing fouling.
- 4. Adding $Ca(OC1)_2$ and H_2O_2 did reduce the amount of color, odor, H_2S_3 , and did increase the amount of dissolved oxygen and bacteria killed, but the results obtained do not justify using this chemical treatment.
- 5. In spite of the problems with lowering the fouling index, the RO system performed well. Based on performance data, the membrane did not appear to have fouling problems. This was substantiated by the small amount of dirt removed during the trial cleaning operation. The use of a spiral-wound membrane, which is less sensitive to fouling, is a better choice than the hollow-fiber membrane.
- 6. Although the ion-exchange deionizer performed as well as the RO system, with respect to TDS removal, the amount of regenerating chemicals required would increase in direct proportion to the concentration of TDS in the feed water. Supply of such large quantity of regenerating chemicals will create severe logistic burden to the Navy. Based on performance data, resins fouling did not appear to be a problem.
- 7. The RO system has an added advantage over the ion-exchange deionizer: the RO system is capable of removing other contaminants, such as colloidal particles, organic chemicals, and micro-organisms not removed in the pretreatment components; the system also uses less chemicals.
- 8. Cost of the RO system has been reduced substantially during the last few years. The performance of the RO membranes has also improved. The RO system has been rated as the least cost desalination process and most feasible water treatment system for Diego Garcia.

- 9. The water produced by the water treatment system is too low in TDS and too acidic to be suitable for long-term human consumption. The product water must be blended with undemineralized water to provide water of potable quality.
- 10. The total TTHMs content in the well water and the present water supply appears to be low (below EPAs 100 ppb standard); there is no TTHMs health threat to the personnel. The water treatment systems tested were effective in removing TTHMs.

RECOMMENDATIONS

- 1. The RO water treatment system is recommended for use at Diego Garcia for treating brackish water as supplemental to the existing fresh water supply.
- 2. Should there be no adequate source and quantity of fresh/brackish water on the island available, seawater desalting RO system is recommended for use.
- 3. When RO water treatment system is in use, the addition of power recovery system to reduce power consumption is recommended.
- 4. One pretreatment alternative that should be strongly considered is an ultrafiltration (UF) membrane process. The UF membrane module is constructed with the same precision as RO modules. The UF module does not reject salts, but removes practically all organics over 1,000 molecular weight (e.g., polysaccharides, virus, bacteria, and colloids).
- 5. The waste stream from a demineralization plant processing brackish water in the 1,000 to 3,000 ppm TDS range will usually constitute from 10 to 30% of the water fed and will contain from 5,000 to 10,000 ppm TDS. At Diego Garcia, the RO system waste stream contained about 2,000 ppm TDS (from 550 ppm water fed). An appropriate disposal option must be provided for the waste; such as discharged to lagoon, ocean and/or injected to ground.

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Cyclo Chemical Corp., Cyclose ion exchange celluloses, Santa Fe Springs, Calif.

Rohm and Haas., Deionization with amberlite MB-A, Philadelphia, Penn.

Appendix A

WATER TREATMENT SYSTEM OPERATION MANUALS

- 1. Cat. No. 8802-81, HI-FLO DEPTH & CULLAR FILTER MEDIA FILTERS, Model 20, Installation Instructions.
- 2. Cat. No. 8805-87, DT-60 FEEDERS, Installation Instructions.
- 3. Cat. No. 8176-86, CULLIGAN A SERIES TWO COLUMN INDUSTRIAL DEIONIZER.
- 4. Culligan. INSTALLATION AND OPERATING INSTRUCTIONS FOR A KD-20 REVERSE OSMOSIS SYSTEM.
- 5. Gould Pump, Model 3333, INSTALLATION, OPERATION AND MAINTENANCE.

Installation, Operation, and Maintenance Manuals for:

- 1. HI-FLO DEPTH FILTER CULLIGAN MODEL HD-20
- 2. CULAR CARBON FILTER CULLIGAN MODEL HR-24
- 3. POLYELECTROLYTE FEEDER CULLIGAN MODEL CT-10
- 4. TWC COLUMN INDUSTRIAL DEIONIZER CULLIGAN MODEL
- 5. REVERSE OSMOSIS UNIT CULLIGAN MODEL KD-20
- 6. MULTI STAGE CENTRIFUGAL PUMP GOULD MODEL 3333

Appendix B

WELL WATER QUALITY

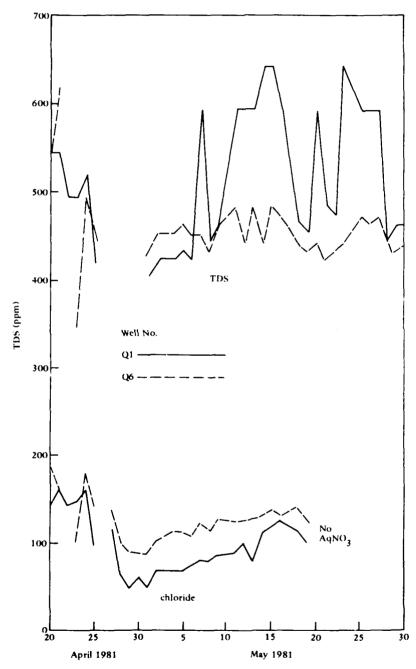


Figure B-1. TDS and chloride concentration correlation.

Table B-1. Well Water Quality, Characteristics, and TDS-Chloride Concentration Correlation

| | D.0. (ppm) | 5.5 | | 3.2 | 0.9 | 2.8 | | 1.8 | 1.6 | 3.6 | 3.6 | 9.9 | 4.8 | 8.9 |
|---------------|------------------------------------|---------|---------|---------|---------|---------|--------------|---------|---------|---------|---------|---------|---------|---------|
| | DS/TDS (ppm) | 550 | | 900 | 625 | 400 | | 909 | 625 | 009 | 575 | 375 | 375 | 500 |
| | Ca Hardness (ppm) | 89 | | 44 | 38 | 100 | | 100 | 72 | 09 | 84 | 84 | 84 | 26 |
| | lotal Hardness (ppm) | 240 | | 500 | 170 | 200 | | 264 | 246 | 186 | 500 | 186 | 500 | 192 |
| lity | Chloride (ppm) | 168 | | 120 | 172 | 118 | | 228 | 233 | 175 | 125 | 72 | 09 | 199 |
| Water Quality | Alkalinity Chloride (ppm) (ppm) | 850 | | 850 | 700 | 650 | | 650 | 750 | 550 | 200 | 009 | 700 | 550 |
| | Turbidity (JTU) | 4 | | က | က | Z. | conductivity | 8 | m | 4 | m | 4 | m | 5 |
| | Color | 20 | | 50 | 20 | 30 | high | 20 | 30 | 10 | 30 | 20 | 20 | 20 |
| | Hd | 6.8 | | 7.2 | 7.5 | 7.0 | due to | 6.8 | 7.0 | 7.2 | 7.0 | 7.0 | 6.8 | 7.5 |
| ; | Sampling Dates | 4/21/81 | 4/21/81 | 4/21/81 | 4/21/81 | 4/22/81 | secured d | 4/22/81 | 4/22/81 | 4/22/81 | 4/22/81 | 4/22/81 | 4/22/81 | 4/22/81 |
| : | Number | 0-1 | 0-3 | 0-4 | 9-0 | C-1 | C-2 | C-3 | C-4 | C-5 | 9-0 | C-7 | ۵-2 | 6-3 |

(continued)

Table 8-1. Continued

| | | | | | Water Quality | lity | | | | |
|----------------|-------------------|-----|-------|--------------------|---------------------|-------------------|----------------------------|-------------------------|-----------------|---------------|
| Well Number | Sampling Dates | Hd | Color | Turbidity (JTU) | Alkalinity (ppm) | Chloride (ppm) | Total Hardness (ppm) | Ca Hardness (ppm) | DS/TDS (ppm) | D.O. (ppm) |
| 0-1 | 4/29/81 | 7.2 | 10 | 5 | 007 | 55 | 160 | 65 | 100 | 5.8 |
| 0-4 | 4/29/81 | 7.0 | 30 | 10 | 920 | 72 | 155 | 100 | 125 | 4.0 |
| 9-0 | 4/29/81 | 7.5 | 50 | m | 200 | 96 | 152 | 95 | 150 | 9.9 |
| C-1 | 4/30/81 | 7.2 | 30 | ъ | 009 | 77 | 200 | 124 | 165 | 4.8 |
| C-3 | 4/30/81 | 7.7 | 30 | m | 929 | 72 | 224 | 148 | 165 | 5.6 |
| C-4 | 4/30/81 | 7.0 | 50 | m | 200 | 84 | 216 | 132 | 180 | 4.8 |
| C- 5 | 4/30/81 | 7.2 | 10 | 2 | 700 | 77 | 188 | 112 | 160 | 5.4 |
| 9-0 | 4/30/81 | 7.5 | 10 | 2 | 750 | 77 | 164 | 104 | 160 | 9.9 |
| 2-3 | 4/30/81 | 7.0 | 10 | 7 | 650 | - 69 | 172 | 132 | 150 | 3.0 |
| 8-J | 4/30/81 | 7.0 | 10 | 2 | 750 | 72 | 212 | 148 | 160 | 5.0 |
| 6-3 | 4/30/81 | 7.0 | 10 | 2 | 550 | 48 | 156 | 108 | 130 | 3.4 |
| P1-13 | 4/30/81 | 7.5 | 10 | | 200 | 43 | 140 | 88 | 100 | 6.2 |
| | | | | | | | | | | } |

(continued)

Table B-1. Continued

| : | | | | | Water Quality | lity | | | | |
|--------|-------------------|-----|-------|--------------------|---------------------|-------------------|----------------------------|-------------------------|-----------------|---------------|
| Number | Sampling Dates | 풥 | Color | Turbidity (JTU) | Alkalinity (ppm) | Chloride (ppm) | iotai Hardness (ppm) | Ca Hardness (ppm) | DS/TDS (ppm) | 0.0. (ppm) |
| 0-1 | 5/5/81 | 7.0 | 20 | 3 | 450 | 72 | 168 | 120 | 440 | 4.8 |
| 0-4 | 5/5/81 | 7.0 | 50 | Z. | 450 | 82 | 168 | 100 | 440 | 3.4 |
| 9-0 | 5/5/81 | 7.7 | 30 | S. | 200 | 120 | 160 | 08 | 470 | 0.9 |
| C-1 | 5/6/81 | 7.0 | 20 | m | 650 | 84 | 172 | 108 | 440 | 3.2 |
| C-2 | 5/6/81 | 7.0 | 20 | -1 | 550 | 180 | 220 | 120 | 700 | 4.4 |
| C-3 | 2/6/81 | 7.0 | 20 | - | 009 | 456 | 280 | 100 | 1,400 | 1.4 |
| C-4 | 5/6/81 | 7.0 | 10 | - | 200 | 252 | 240 | 96 | 1,000 | 1.4 |
| C-5 | 18/9/5 | 7.2 | 10 | 7 | 200 | 113 | 180 | 100 | 470 | 5.0 |
| 9-0 | 2/6/81 | 7.0 | 10 | | 200 | 113 | 200 | 120 | 009 | 4.2 |
| C-7 | 5/6/81 | 7.0 | 10 | | 200 | 28 | 180 | 120 | 420 | 4.4 |
| 8-3 | 5/6/81 | 7.0 | 10 | - | 009 | 09 | 200 | 132 | 440 | 4.6 |
| 6-3 | 5/6/81 | 7.2 | 10 | 7 | 450 | 144 | 180 | 64 | 490 | 5.0 |
| P1-13 | 5/6/81 | 7.5 | 10 | 0 | 350 | 24 | 120 | 80 | 300 | 6.6 |

(continued)

Table 8-1. Continued

| | D.O. (ppm) | 5.4 | 5.8 | 6.4 | 5.4 | 1.6 | 8.4 | 6.2 | 2.0 | 4.8 | 2.4 | 5.6 | 3.8 | 5.0 | 3.6 | 3.8 | 6.2 | (pər |
|---------------|----------------------------|--------|--------|--------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|-------------|
| | DS/TDS (ppm) | 009 | 360 | 460 | 200 | 909 | 470 | 480 | 450 | 200 | 800 | 009 | 009 | 400 | 430 | 009 | 250 | (continued) |
| | Ca Hardness (ppm) | 128 | 104 | 128 | 120 | 148 | 112 | 100 | 120 | 132 | 124 | 128 | 104 | 116 | 140 | 100 | 96 | |
| | Total Hardness (ppm) | 200 | 148 | 192 | 156 | 184 | 144 | 144 | 180 | 188 | 240 | 180 | 200 | 172 | 200 | 180 | 120 | |
| lity | Chloride (ppm) | 98 | 09 | 108 | 108 | 120 | 91 | 132 | 84 | 168 | 163 | 156 | 120 | 09 | 09 | 168 | 53 | |
| Water Quality | Alkalinity (ppm) | 200 | 650 | 009 | 550 | 059 | 200 | 200 | 009 | 650 | 650 | 650 | 009 | 650 | 450 | 200 | 400 | |
| | Turbidity (JTU) | 7 | 2 | m | т | m | က | က | က | က | 2 | 2 | ĸ | 2 | 2 | 2 | 2 | |
| | Color | 10 | 10 | 20 | 30 | 30 | 30 | 30 | 30 | 30 | 10 | 30 | 30 | 10 | 10 | 70 | 20 | |
| | Ha | 7.0 | 7.7 | 7.2 | 7.2 | 7.0 | 7.0 | 7.7 | 7.0 | 7.2 | 7.0 | 7.2 | 7.0 | 7.2 | 7.0 | 7.7 | 7.7 | |
| | Sampling Dates | 5/7/81 | 5/7/81 | 5/8/81 | 5/12/81 | 5/12/81 | 5/12/81 | 5/12/81 | 5/13/81 | 5/13/81 | 5/13/81 | 5/13/81 | 5/13/81 | 5/13/81 | 5/13/81 | 5/13/81 | 5/13/81 | |
| | Well | 0-1 | 6-0 | 9-0 | 0-1 | 0-3 | 4-0 | 9-0 | C-1 | C-2 | C-4 | C-5 | 9-0 | C-7 | 8-5 | 6-0 | P1-13 | |

B-6

Table B-1. Continued

| | D.O. (ppm) | 5.4 | 1.4 | 5.5 | 6.0 | 1.6 | 1.4 | 3.8 | 4.2 | 3.2 | 3.8 | 3.0 | 6.0 |
|---------------|----------------------------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|
| | DS/TDS (ppm) | 460 | 200 | 420 | 440 | 400 | 700 | 470 | 200 | 380 | 400 | 009 | 240 |
| | Ca Hardness (ppm) | 108 | 136 | 84 | 64 | 4 | 70 | 100 | 100 | 108 | 128 | 80 | 80 |
| | Total Hardness (ppm) | 180 | 240 | 144 | 156 | 176 | 200 | 160 | 196 | 172 | 192 | 172 | 120 |
| lity | Chloride (ppm) | 108 | 139 | 91 | 137 | | | | - | | | | |
| Water Quality | Alkalinity (ppm) | 009 | 750 | 200 | 200 | 650 | 009 | 450 | 009 | 200 | 009 | 450 | 300 |
| | Turbidity (JTU) | 1 | က | 2 | S | m | က | က | m | m | m | m | 2 |
| | Color | 20 | 30 | 50 | 30 | 30 | 30 | 30 | 30 | 30 | 30 | 30 | 30 |
| | Ŧ | 7.2 | 6.8 | 7.2 | 7.7 | 7.2 | 7.2 | 7.5 | 7.2 | 7.2 | 7.0 | 7.2 | 7.5 |
| | Sampling Dates | 5/19/81 | 5/19/81 | 5/19/81 | 5/19/81 | 5/20/81 | 5/20/81 | 5/20/81 | 5/20/81 | 5/20/81 | 5/20/81 | 5/20/81 | 5/20/81 |
| | Well | 0-1 | 0-3 | 4-0 | 9-0 | C-1 | C-4 | C-5 | 9-0 | C-7 | 8-5 | 6-3 | P1-13 |

(continued)

Table B-1. Continued

| | | | | | Water Quality | lity | | | | |
|----------------|-------------------|-----|-------|--------------------|---------------------|-------------------|----------------------------|-------------------------|-----------------|---------------|
| Well Number | Sampling Dates | 푎 | Color | Turbidity (JTU) | Alkalinity (ppm) | Chloride (ppm) | Total Hardness (ppm) | Ca Hardness (ppm) | DS/TDS (ppm) | D.0. (ppm) |
| | 5/26/81 | 7.2 | 20 | 2 | 750 | | 200 | 120 | 009 | 9.9 |
| | 5/26/81 | 6.8 | 30 | m | 750 | | 268 | 148 | 750 | 9.0 |
| | 5/26/81 | 7.2 | 20 | m | 009 | | 192 | 116 | 460 | 4.6 |
| | 5/26/81 | 7.7 | 30 | m | 200 | | 176 | 92 | 470 | 6.0 |
| | 5/26/81 | 7.0 | 30 | m | 009 | | 184 | 80 | 420 | 6.4 |
| | 5/27/81 | 7.2 | 35 | m | 009 | | 220 | 09 | 200 | 5.4 |
| | 5/27/81 | 7.2 | 20 | m | 250 | | 168 | 100 | 430 | 5.0 |
| | 5/27/81 | 7.5 | 35 | S | 009 | | 216 | 100 | 490 | 5.4 |
| C2 | 5/27/81 | 7.2 | 20 | 7 | 550 | | 180 | 104 | 380 | 6.4 |
| <u> </u> | 5/27/81 | 7.2 | 30 | 2 | 009 | | 192 | 128 | 400 | 8.9 |
| | 5/27/81 | 7.5 | 30 | ഹ | 200 | | 172 | 84 | 009 | 3.8 |
| P1-13 | 5/27/81 | 7.7 | 30 | ~ | 200 | | 116 | 88 | 240 | 8.9 |

Appendix C

PRETREATMENT TEST RESULTS

Table C-1. pH adjustment

Experiment 1. July 1, 1981. Well water alkalinity = 600 ppm as $CaCO_3$ pH initial = 7.05. Well water volume = 1,000 ml.

Experiment 2. July 2,1981. Well water alkalinity = 620 ppm as $CaCO_3$ pH initial = 7.6. Well water volume = 1,000 ml.

Acid strength used in Experiment 1 and 2, 1/10 of 20°Be HC1.

Experimental results:

| Experime | ent 1 | Experime | nt 2 |
|---|---|---|--|
| Acid Volume (m1) | ph Value | Acid Volume (ml) | pH Value |
| 0 0.5 1.0 1.5 2.0 2.5 3.0 3.5 4.0 4.5 5.0 5.5 6.0 6.5 7.0 7.5 8.0 | 7.05 6.7 6.5 6.2 6.15 6.0 5.9 5.8 5.65 5.35 5.1 4.6 3.4 3.2 3.2 | 0 0.5 1.0 1.5 2.0 2.5 3.0 3.5 4.0 4.5 4.75 5.0 5.25 5.5 6.0 6.5 7.0 7.5 8.0 | 7.6 7.25 7.0 6.7 6.6 6.35 6.2 6.0 5.8 5.75 5.65 5.4 4.6 3.45 3.45 3.45 3.2 |

Flow rate further reduces down to 30 ml/mir (1300 H) 1350 - added 3 gm of HTH to about 60 gal of water; 1352 - residual Cl₂ = 6 ppm Added approx
10 ppm of
Ca(OC1)
Measured fmmediacely. Obtained 2 ppm
Cl. residual.
Reduce flow
rare from 50
to 35 ml/min.
Partial plug-up. $H_2^{0_2}$ added $H_2^{0_2}$ added $H_2^{0_2}$ added Remarks Residual Chlorine 3.0 0.0 0.0 0.0 Flow Rate (ml/min) 32.5 8 38 38 35 36 28 23 Bacteria 900 DINI DINI 200 100 909 200 Table C-2. Test Results of Slow Sand Filtration 7.7 7.5 8.0 7.7 1.1 8.0 7.7 7.7 7.7 Ħ Alkaliníty 580 580 580 580 580 630 630 480 680 630 580 680 530 630 580 580 580 Chloride 66 66 108 108 108 106 106 106 108 108 104 127 115 Dissolved Oxygen 11.2 6.6 6.6 7.6 7.2 7.8 4.9 Calcium Hardness 120 100 120 120 120 96 96 100 100 100 112 96 112 96 112 128 9 112 112 210 168 152 172 156 168 196 164 320 180 172 172 172 176 164 164 172 172 75 176 168 067 087 067 470 7,60 480 Turbidity m0 m ~ 0 0 7 0 m 0 m 0 m 0 ma ma Color 25 25 2 8 Influent Effluent Influent Effluent Influent Effluent Influent Effluent Influent Effluent Influent Effluent Influent Effluent Effluent Influent Influent Effluent 1300 0900 7-20-81 0900 7-20-81 0900 1330 0900 0830 1230 0900 7-23-81 Hour and Date

.

Cont inued

Table C-2. Continued.

| ω e − | TUS (ppm | Turbidity (ppm) | E gr | | Total Hardness | Calcium Hardness | Dissolved Oxygen | Chloride | Chloride Alkalinity | Ηď | Bacteria | Flow Rate (ml/min) | Residual Chlorine | Remarks |
|--------------------|-------------|-----------------|------|---|-------------------|---------------------|---------------------|----------|---------------------|-----|----------|--------------------------|----------------------|--|
| 136 120 164 104 | - | 176 | 176 | - | 220 | 0.4 | 6.6 | 115 | 630 530 | 8.0 | | 25 | | Experimental set-up was transferred to another plant |
| 176 120 156 104 | | 176 | 176 | | 120 | 0.1 | 6.6 | 108 | 680 580 | 8.0 | | 22 | | |
| 168 120 152 104 | | 168 | 168 | | 120 | 0.1 | 6.2 5.6 | 108 | 680 580 | 8.0 | | 17 | | |
| 168 120 152 92 | | 168 | 168 | | 120 | | 8.0 | 97 | 630 580 | 8.0 | | 16 | | |
| 168 120 152 100 | | 168 | 168 | | 10,12 | 0.0 | 9.9 | 92 | 530 | 7.9 | | 115 | | |
| 164 104 104 104 | | 164 | 164 | - | 01 & | 40 | 6.4 5.6 | 95 | 630 | 8.0 | | 15 | | |

Table C-3. Fouling Index

| Date | Multi-Media Filter Effluent |
|----------|--------------------------------|
| 05-01-81 | 3.10 |
| 05-02-81 | 2.70 |
| 05-07-81 | 6.10 |
| 05-08-81 | 6.10 |
| 05-09-81 | 5.70 |
| 05-14-81 | 4.50 |
| 05-15-81 | 4.50 |
| 05-16-81 | 4.80 |
| 05-21-81 | 4.80 |
| 05-22-81 | 4.80 |
| 05-23-81 | 5.00 |
| 07-15-81 | 4.20 |
| 07-24-81 | 7.46 |
| 07-25-81 | 6.71 |
| 07-28-81 | 5.69 |
| 07-29-81 | 6.14 |
| 07-30-81 | 6.59 |
| 07-31-81 | 7.50 |
| 08-01-81 | 6.31 |

Table C-4. Chemical Additions

Experiment No.1. Hydrogen peroxide added to raw well water

| Sample | H ₂ O ₂ Dosage (ppm) | Color | Sulfide | DO | Odor |
|--------|---|-------|---------|------|--------|
| Blank | 0 | 30 | 0.3 | 4.2 | Yes |
| 1 | 2.5 | 30 | 0.1 | 5.4 | Slight |
| 2 | 5.0 | 30 | 0 | 6.0 | No |
| 3 | 7.5 | 25 | 1 0 | 8.5 | No |
| 4 | 10.0 | 25 | 1 0 | 8.6 | No |
| 5 | 12.5 | 20 | 0 | 11.0 | No |
| 6 | 15.0 | 20 | Ō | 11.0 | No |

Experiment No. 2. Calcium hypochlorite and hydrogen peroxide added to raw well water

| Sample No. | Ca(CO1 ₂) (ppm) | H ₂ O ₂ (ppm) | 0dor | Color | C1 ₂ Residual | Sulfide | Bacteria Count |
|------------------|--------------------------------|--|-----------------------|----------------------------|-----------------------------|-----------------|---------------------|
| B 1 2 3 | 0 2.5 5.0 | 0 0 0 2.5 5.0 | Yes No No No | 30 30 30 30 30 | 0 0 0 | 0.4 0.1 0 | 45 30 25 3 |

Appendix D

ION-EXCHANGE DEIONIZER'S OPERATION DATA SUMMARY AND PERFORMANCE DATA SHEETS

(continued)

NaOH (gal) 5555 Chemicais HCl (gal) 626626 Percentage of Input Ion-Exchange Deionizer Operation Data Summary 87.9 87.3 87.3 87.3 86.2 86.2 100 100 100 84.5 84.5 84.5 86.3 100 86.4 86.4 86.4 86.4 86.7 70.8 85.4 70.8 69.6 85.4 70.8 Product Performance Flow Rate (gpm) for this period 8.9 8.8 8.8 8.2 7.2 7.2 7.2 6.5 6.5 7.6 10.5 4 21.6 21.6 18.4 16.5 16.1 18.2 13.4 test) Regeneration (hrs) Wo operating data available at start of reading 17 20.5 17 16 20.5 17 Table D-1. Total^a (hrs) (Meter 10,885 10,795 10,320 6,900 6,730 9,360 4,760 1,150 1,290 8,630 18,750 20,270 16,390 17,470 16,470 12,930 Product (gal) 8,220 3-15-81 3-16-81 3-17-81 3-18-81 3-20-81 3-21-81 3-22-81 3-25-81 3-25-81 3-26-81 3-31-81 4-1-81 4-1-81 4-1-81 4-19-81 4-20-81 4-21-81 4-22-81 4-27-81 4-28-81 Date

Table D-1. Continued

| Product Performance |
|---------------------|
| Flow Rate (gpm) |
| 16.3 |
| 16.5 |
| 16.4 |
| 14.9 |
| 18.2 |
| 16.5 |
| 17.3 |
| 17.5 |
| 18 |
| 20 |
| 17. |
| 17 |
| |
| 13.9 |
| |
| |
| |
| |
| |
| |
| |

 a Time period = 0800 - 0800.

^bBased on 1,500 gallons of water for regeneration.

Regeneration cycle was started accidentally.

dAdded booster pump to increase flow rate to the system.

^eIncreased flow rate to maximum.

Table D-2. Summary of Water Analyses of Feed to and Water Product from a Ion-Exchange Deionizer

| Turbidity (JTU) | Out | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | | | |
|------------------------------|-----|--------|--------------------|---------|---------|---------|---------|--------|--------|---------|---------|---------|---------|---------|----------|---------|----------|----------|-----|------|---------|
| Turb (3 | In | 7 | ი ~ | - | 7 | ~ | m | ო | 2 | m | 7 | 7 | 7 | ~ | m | m | m | 7 | | | |
| Color | 0ut | 0 | <u> </u> | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | | | |
| ၀၁ | υI | 02 | 52 52 | 20 | 20 | 20 | ಜ | 20 | 20 | ဣ | 9 | 9 | 9 | 8 | 8 | ဓ္က | 50 | 30 | | | |
| Hd | 0ut | | v. v. | | | | | | | | | | | | | | | | | | |
| d | In | 7.5 | 7.5 | 7.6 | 7.5 | 7.5 | 7.5 | 7.5 | 7.7 | 7.7 | 7.7 | 7.7 | 7.7 | 7.2 | 7.5 | 7.7 | 7.7 | 7.7 | | | |
| inity m) | 0ut | 150 | 150 | 300 | 150 | 150 | 200 | 100 | 200 | 200 | 150 | 200 | 250 | 100 | 200 | 150 | 200 | 150 | | | |
| Alkalinity (ppm) | In | 650 | /50 650 | 650 | 700 | 650 | 550 | 450 | 200 | 220 | 650 | 200 | 009 | 009 | 450 | 009 | 009 | 220 | | | - |
| ride m) | Out | 7 | 76 | 20 | 64 | 72 | 78 | 19 | 19 | 24 | 24 | 10 | 12 | ; | ; | ; | ; | ; | of | ride | ent |
| Chloride (ppm) | In | 115 | 124 | 110 | 105 | 86 | 9/ | 96 | 84 | 98 | 96 | 96 | 84 | ; | : | ! | ; | ; | 0ut | chlo | reagent |
| ium ness m) | 0ut | ő | 00 | 01 | 80 | 15 | 15 | 0 | 0 | 4 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | | | |
| Calcium Hardness (ppm) | In | 248 | 115 | 135 | 107 | 123 | 115 | 104 | 100 | 104 | 120 | 108 | 100 | 108 | 96 | 96 | 100 | 84 | | | |
| al ness (n | Out | Õ | 12 | 0 | 12 | 15 | 10 | 0 | 21 | 16 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | | | |
| Total Hardness (ppm) | In | 372 | 198 | 174 | 185 | 197 | 178 | 168 | 172 | 180 | 180 | 172 | 169 | 172 | 172 | 168 | 180 | 180 | | | |
| ; <u>(</u> | Out | 12 | 77 | 42 | 91 | 52 | 12 | 6 | _ | 10 | 10 | 9 | 6 | <u></u> | <u>о</u> | 4 | ∞ | ∞ | | | |
| 70S (ppm) | In | 009 | 425 425 | 400 | 225g | 175 | 75ª | 430 | 430 | 460 | 400 | 460 | 370 | 300 | 370 | 400 | 410 | 400 | | | |
| Date | | 4-3-81 | 4-20-81 4-21-81 | 4-22-81 | 4-27-82 | 4-28-81 | 4-29-81 | 5-4-81 | 5-5-81 | 5-11-81 | 5-12-81 | 5-13-81 | 5-18-81 | 5-19-82 | 5-20-81 | 5-25-81 | 5-26-81 | 5-27-81 | | | |
| i | | ŀ | | | | | | | | | | | | | | | | | | | |

^aMeter was out of order on these dates.

Appendix E

RO DEMINERALIZER OPERATIONAL AND PERFORMANCE SUMMARY

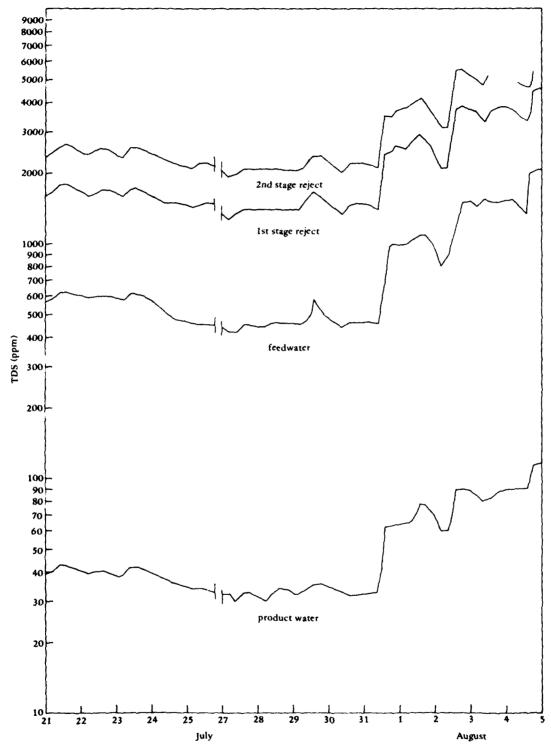


Figure E-1. Correlation of TDS in product water and feedwater; first- and second-stage rejects.

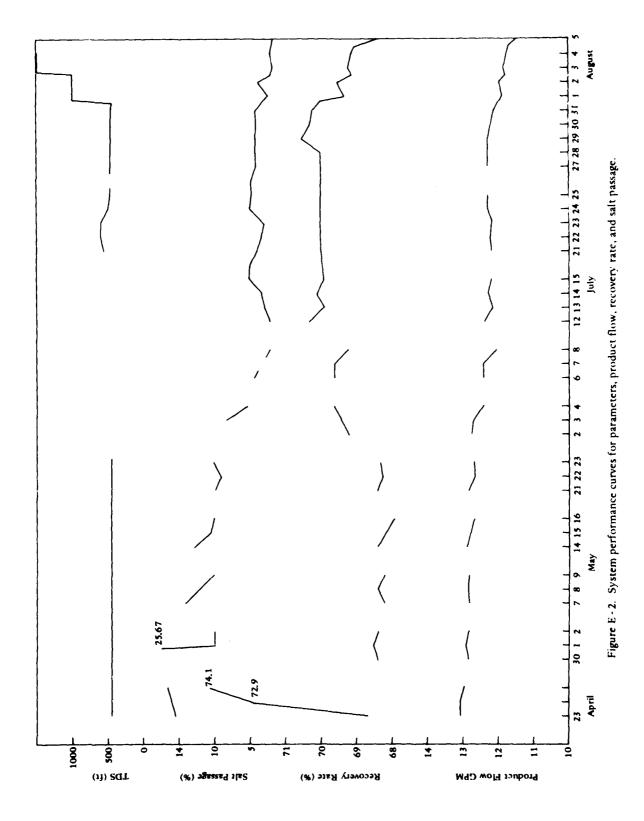


Table E-1. Demineralizer Operational Summary

| | Time | Fì | ows | | Inlet P | ressure | Perfor | mance |
|--------------------|---------------------|------------------------------------|---------------------------|-------------------------|--------------------------|---------------------------|-------------------------|------------------------|
| Date | on Line (hrs) | Total Amount of Water (gals) | Product Water (gpm) | Waste Water (gpm) | First Stage (psig) | Second Stage (psig) | Recovery Rate (%) | Salt Passage (%) |
| 4-23-81 | 4 | 4,800 | 13.8 | 6.0 | 400 | 370 | 69.7 | 12.75 |
| 4-24-81 | 24 | 27,900 | 13.8 | 5.1 | 400 | 370 | 72.9 | 12.97 |
| 4-25-81 | 24 | 26,700 | 13.7 | 4.7 | 400 | 370 | 74.1 | 13.32 _a |
| 4-30-81 | 8 | 9,600 | 13.6 | 6.0 | 400 | 370 | 69.4 | _ |
| 5-1-81 | 24 | 25,500 | 13.7 | 6.0 | 400 | 370 | 69.5 | 9.90 |
| 5-2-81 | 24 | 28,400 | 13.6 | 6.0 | 400 | 370 | 69.4 | 9.92 |
| 5-7-81 | 8 | 9,500 | 13.5 | 6.0 | 400 | 370 | 69.2 | 12.03 |
| 5-8-81 | 24 | 23,000 | 13.6 | 6.0 | 400 | 370 | 69.4 | 10.82 |
| 5-9-81 | 24 | 27,500 | 13.5 | 6.0 | 400 | 370 | 69.2 | 10.0 |
| 5-14-81 | 8 | 8,300 | 13.6 | 6.0 | 400 | 370 | 69.4 | 11.4 |
| 3-15-81 | 24 | 28,300 | 13.4 | 6.0 | 400 | 370 | 69.1 | 10.23 |
| 5-16-81 | 24 8 | 27,500 | 13.3 13.6 | 6.0 | 400 | 370 | 68.9 | 10.05 |
| 5-21-81 | 24 | 5,350 | 13.6 | 6.0 5.9 | 400 400 | 370 370 | 69.4 69.3 | 9.86 |
| 5-22-81 5-23-81 | 24 | 32,050 27,500 | 13.3 | 5.9 | 400 | 370 | 69.3 | 9.45 10.0 |
| 7-2-81 | 8 | 9,200 | 13.4 | 5.7 | 400 | 370 | 70.2 | 10.0 |
| 7-3-81 | 24 | 17,100 | 13.4 | 5.6 | 400 | 370 | 70.2 | 9.1 |
| 7-4-81 | 24 | 33,500 | 13.0 | 5.4 | 400 | 370 | 70.6 | 7.61 |
| 7-6-81 | 24 | 26,800 | 13.0 | 5.4 | 400 | 38=70 | 70.6 | 7.09 |
| 7-7-81 | 14 | 15,500 | 13.0 | 5.4 | 400 | 370 | 70.6 | |
| 8-8-81 | 36 | 34,900 | 12.5 | 5.3 | 400 | 370 | 70.2 | 6.0 |
| 7-12-81 | 10 | 5,370 | 12.9 | 5.2 | 400 | 372 | 70.2 | 5. 97 |
| 7-13-81 | 18 | 20,870 | 12.7 | 5.2 | 400 | 375 | 71.3 | 6.45 |
| 7-14-81 | 24 | 23,900 | 12.8 | 5.2 | 400 | 372 | 71.1 | 6.61 |
| 7-15-81 | 24 | 24,970 | 12.7 | 5.2 | 400 | 370 | 70.9 | 7.51 |
| 7-21-81 | 24 | 25,780 | 12.7 | 5.2 | 400 | 37 <i>2</i> | 71.0 | 6.8 |
| 7-22-81 | 23 | 24,840 | 12.8 | 5.2 | 400 | 375 | 71.0 | 6.5 |
| 7-23-81 | 14.5 | 15,575 | 12.7 | 5.2 | 400 | 375 | 71.0 | 7.5 |
| 7-24-81 | 11 | 11,880 | 12.8 | 5.2 | 400 | 375 | 71.0 | 7.4 |
| 7-25-81 | 23 | 24,840 | 12.8 | 5.2 | 400 | 375 | 71.0 | 7.1 |
| 7-27-8. | 24 | 25,920 | 12.8 | 5.2 | 400 | 375 | 71.0 | 7.2 |
| 7-28-81 | 21 | 22,680 | 12.8 | 5.2 | 400 | 375 | 71.0 | 7.2 |
| 7-29-81 | 24 | 25,920 | 12.8 | 5.1 | 400 | 375 | 71.5 | 7.2 |
| 7-30-81 | 20 | 21,360 | 12.7 | 5.1 | 400 | 375 | 71.3 | 7.1 |
| 7-31-81 | 14 _a | 14,870 | 12.6 | 5.1 | 400 | 375 | 71.2 | 7.1 |
| 1 | 19a | 9,450 | 12.4 | 5.1 | 400 | 375 | 71.0 | 6.7 |
| 8-1-81 | 23 | 24,150 | 12.3 | 5.1 | 400 | 375 | 70.3 | 6.2 |
| 8-2-81 | 18 _b | 19,010 5,220 | 12.4 12.2 | 5.2 5.2 | 400 400 | 375 375 | 70.5 70.1 | 5.8 6.0 |
| 8-3-81 | 20 | 20,940 | 12.25 | 5.2 | 400 | 375 | 70.1 | 5.8 |
| 8-4-81 | 16 5 | 17,225 | 12.2 | 5.2 | 400 | 375 | 70.2 | 6.0 |
| 0 7 01 | 16.5 6.5 | 6,750 | 12.1 | 5.2 | 400 | 375 | 70.0 | 5.8 |
| 8-5-81 | 9.5 | 9,180 | 11.8 | 5.2 | 400 | 375 | 69.4 | 5.7 |
| J J U1 | 9 14 | 14,700 | 12.3 | 5.2 | 400 | 375 | 70.3 | 8.0 |

 $^{^{\}rm a}$ TDS increased to 1,000 ppm in feed stream.

 $^{^{\}mathrm{b}}\mathrm{TDS}$ increased to 1,500 ppm in feed stream.

 $^{^{\}text{C}}\text{TDS}$ increased to 2,000 ppm in feed stream.

dReturn to normal TDS of 450 ppm.

TI Mixed Media Filter 7.46 6.71 6.59 7.5 6.31 Turbidity (JTU) Out 0 0 0 33333335 of Feed Water and Product Water from the RO Demineralizer Out 00 Color 22222222222 828 8 3 യയയ 5.5 Out 50 សំសំសំសំសំសំសំសំសំសំសំសំសំសំ പ്പ്പ്പ് אי אי 돒 7.7 Ë Alkalinity (ppm) Out 200 200 200 200 200 200 200 200 350 350 150 130 180 230 230 230 180 230 190 700 650 650 650 700 700 700 700 800 800 800 600 600 600 600 700 630 530 530 680 530 630 680 580 In I 34 16 16 18 18 23 23 23 23 35 Out 52 52 52 52 36 36 29 29 29 29 29 Chloride (ppm) 1110 1110 1110 1108 1108 87 88 88 88 83 83 92 347 437 Ľ Table E-2. Water Analyses Summary £13121200040004 0 0 4 4 4 0 4 8 Out Calcium Hardness (ppm) 124 124 132 132 132 120 120 120 120 128 Ľ 20002220409900990 Out Total Hardness (ppm) 172 176 176 172 172 164 172 199 199 199 199 199 199 199 199 199 176 188 172 180 228 224 Ľ Out TÜS (PPM) 460 424 444 455 455 11,000 11,500 11,550 2,100 2,100 575 625 600 625 드 4-23-81 4-24-81 4-24-81 5-7-81 5-7-81 5-7-81 5-14-81 5-13-81 7-2-Date

 $^{ extsf{a}}$ Operator not available for simple analyses during 2 thru $extsf{8}$ July 1981.

continued

Table E-3. Preliminary design of a 300,000-gpd RO system.

| FEED D | 13 H | | S ASSUNE ARS OLD GFF OF | D TO 1 UITH 4 | MAS ASSUMED TO BE 82.0 F. YEARS OLD UITH A FLUX DECLINE SLOPE OF 00 GPF OR 208.33 GPM AT 70.0 RECOVENTA | INE SLOP | E GF -0.026 EKI. | 97 | | | | | | 208.3 gpm |
|------------------|--|--------------------------|-------------------------------|--------------------------|---|---|--|---------|----------|----------|--------|-------------|------------|-----------|
| TOTAL FEED PU | TOTAL NUMBER OF ELEMENTS IN THIS SYSTEM IS FEED PRESSURE = 428.7 PSI CONC. PRESSUR | ELEMENTS IN 428.7 PSI | IN THIS | SYSTE! | IS SYSTEM IS 36. CONC. PRESSURE = | 385.4 851 | 1-4 | | | | | 4 | | |
| BANK | r 4. | EED FL | : 000 | COM | COMC, FLOW | BETA | CONC. PRES | RES | | | | | 7 _ | |
| C4 | 101AL 277.78 134.39 | 45 - 5 | - 1 | 67.439 67.62 67.62 | 33.60 34.61 | | F51 (.15 406.08 1.12 385.37 278 gpm | 0 | 2 2 2 | | | | 09.68 mm | د |
| | 1 | FEEL | | | 1 2 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 | | คอลทยิดโย- | 1 | | ! | CONCE | CONCENTRATE | - | |
| ION | #6/L | MEQ/L | *#94 | | ION | 1/9H | MED/L | *#44 | - | 101 | 46/1 | MEG/1 | *#44 | |
| Ş | 52.0 | 2.0 | 129.4 | ·- ·• | i, | 6.0 | 0.0 | e. | | C.A | 204.9 | 10.2 | 509.7 | |
| 9¥ | 24.0 | 2.0 | 9.8.4 | | #6 | 0.4 | 0.0 | ·• • | ٠. | 9₩ | 94.6 | 7. | 387.6 | |
| ₹ | 439.0 | 19.1 | 954.3 | | ĄZ. | 37.4 | 1.6 | 81.4 | | NA | 1640.3 | 71.3 | 3565.8 | |
| ·* | 0.0 | | 0.0 | - | غد | 0.0 | 0.0 | 0.0 | | *. | 0 | o .5 | ٥.٥ | |
| ¥n¥ | 0.0 | 0.0 | 9.0 | | አስት | 0.0 | 0,0 | Ø*•9 | - | 2114 | 0.0 | 0.0 | 0.0 | |
| HC03 | 106.0 | | 86.3 | | HC03 | 9 | 0.1 | υ. υ | - • | HC03 | 404.5 | 0.0 | 331.4 | |
| S04 | 484.0 | 10.1 | 504,2 | | 504 | 5.6 | r., | رة 8 | | 504 | 1914.4 | 30.0 | 1994.2 | |
| ಚ | 420.0 | B • 1 | 591.5 | | ส | ار د د د د د د د د د د د د د د د د د د د | <i>ن</i> | 74.7 | ٠- | ಚ | 1519.2 | 47.0 | 6. 65 | |
| NO.3 | 0.0 | 0.0 | 0.0 | - | N03 | 0 | 0.0 | o.0 | | NO.2 | 0.0 | 0.0 | 0.0 | |
| | 0.0 | 0.0 | 0.0 | ٠. | 14. | 0.0 | 0.0 | 0.0 | - • | . | 0.0 | ٥. ٥ | 0.0 | |
| \$102 | 0.0 | • | 0.0 | | 5162 | 6.0 | ٥. ن | 0.0 | | 2018 | 17.2 | 0.0 | 0.0 | |
| 158 | 1530.0 | 1 1 1 | | | 115 | 104.1 | , | 1 | | 195 | 100 | | | |

Table E-3. Continued.

| ທ | ROBERT BLEVITT |
|-------------------------------------|--|
| 8 | ing. |
| THIS IS ROSYS, FOR VERSION OOS MAUT | THIS CALCULATION WAS MADE BY: ROBERT BLEVIT USING BS40 A ELEMENTS |
| THIS I | THIS CUSING |

10-SEP-81 12:16:20 CALCULATION NUMBER 7275

FEED WATER TEMPERATURE WAS ASSUMED TO BE 82.0 F.

THE ELEMENTS WERE 3.00 YEARS OLD WITH A FLUX DECLINE SLOPE OF -0.026

TOTAL PRODUCT = 300,000 GPD OR 208.33 GPM AT 75.0 RECOVERY,

THE ARRAY WAS 4 2

TOTAL NUMBER OF ELEMENTS IN THIS SYSTEM IS 36.

FEED PRESSURE = 503.6 PSI CONC. PRESSUKE = 460.9 PSI

| FEED | | | FLUB | 8618 | CONC. PRES |
|--------|-------|--------|-------|------|------------|
| TOTAL | | | VESSE | ب. | |
| GPM | | | GFK | | PSI |
| 277.78 | 09.44 | 135.08 | 33.77 | 1.15 | 482,31 |
| 35.08 | | | 34.64 | | |

| 100 | 46/5 | HEQ/L | PFMs | | 10k | 1/9# | MEGUL | PPM+ | | ION | 7/9H | MEQ/L | **** |
|------------|--------|---|---|----------|----------|---------|--------|-------------|------------|--------|--------------|----------|-------|
| 4 3 | 52.0 | 2.6 | 129.4 | | CA | C 4 | 0.1 | 6.€ | | 5 | 205.0 | 10.2 | 509.9 |
| : <u>:</u> | 0.40 | 0.0 | 98.4 | | (2) Æ | 3 | 0.0 | 1 | - | 130 | 44.6 | ээ *\ | 387. |
| ٠ | 0.654 | 1. | 954.3 | | Z. | | 03 , 1 | 8.14. | | # W | 1565.1 | 0.89 | 3405 |
| : | 0.0 | 0.0 | 0.0 | . | ** | 0 | 0.0 | 0.0 | | 242 | 0.0 | 0.0 | ÷ |
| AT. | 0.0 | 0 | 0.0 | - | 415 | 0.0 | 0.0 | 0.0 | | NH4 | 0.0 | 0.0 | > |
| EC 0.3 | 106.0 | , r | 86.9 | | HC03 | 11.2 | 0.0 | 6.5 | ٠. | 4003 | 3 0 10 | 4,3 | 321. |
| 40.5 | 484.0 | 10:1 | 504 | | 504 | 0.5 | . 0 | , o | | 504 | 1922.2 | 40.0 | 2002 |
| | 420.0 | | 10,10 | | . ವ | 93.2 | 2.6 | 131.3 | | ದ | 1405.0 | 39.6 | 1978. |
| 1 D N | 0.0 | 0.0 | 0 7 | | ¥03 | 0.0 | 0.0 | 0.0 | - - | N03 | 0.0 | 0.0 | o |
| <u>.</u> | 9 0 | 0,0 | 0.0 | | . | 0.0 | 0.0 | 0.0 | | L_ | 0.0 | 0.0 | Ġ |
| 5102 | . o | 0.0 | 0.0 | . • | 2018 | 1.5 | 0.0 | o. c | <u></u> . | 5102 | 15.4 | 0.0 | ٥ |
| 50.1 | 1536.0 | 1 | 1 | - ·- | 105 | 1.971 | 1 | 1 1 1 | | 135 | 5599.0 | | ! |

Table E-3. Continued,

| AMPLE DATE: | |
|-------------|--|

10-SEP-81 12:04:06 SERIAL NUMBER: 859

THE IONS WERE ASSUMED TO BE REPORTED AS THE IGNS. SUM OF THE CATIONS IS 5.00 MED/L SUM OF THE ANIONS IS -23.66 MED/L

THE ORIGINAL ANALYSIS DID NOT BALANCE BY -65,08 % A CORRECTION OF 18.45 MED/L WAS ADDED 16 14E NA CONC.

| | | -/*/ 10 47- | | | | | ACT IT TO | POPPE DITE OF BUILDING THE PROPERTY OF THE PRO | 10010 |
|---|---|---|---------------|---|------------|------------|---|--|--------------|
| 10% 10% | CONCENTRATION | NOO | CONCENTRATION | zo | | ION ION | | CONCENTRATION | z |
| | | 43CA | LCULATED | 1 | | | | CALCULATED | |
| | | 46/1 | MED/L | FPM AS CACO3 | | | 1/9H | MED, L | PFM AS CACU3 |
| C. | 52,00 | 52.00 | 2.59 | :29.74 | . . | CA | 52.00 | 2.59 | 129.74 |
| #G | | 24,00 | 1.97 | 89.86 | | 9K | 24.00 | . 97 | 93.68 |
| 44 | | 438.99 | 19.09 | 954,34 | | Œ. | 438.99 | 19.09 | 954,34 |
| HC03 | | 708.00 | 11.60 | 580.14 | | HCG3 | :05.75 | 1.73 | 86.65 |
| 504 | | 00.01 | 0.21 | 16.01 | | 504 | 464.14 | 10.08 | 503,90 |
| נו | | 420.00 | 11.84 | 420.00 11.84 592.22 | | ರ : | 420.00 | 11.84 | 592.22 |
| 105 | 1550.00 | 1657.99 | | | | TINS | 1529.89 | | |
| 1 | 1 | 1 | 1 1 1 1 | 2 5 4 1 5 5 1 2 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 | - | | 1 | 1 | |
| -ALKALINITY | 580.00PPM AS CACOS | H AS CACOS | | 580,14 | | | | | 39.98 |
| -HARDNESS | 224.00PP) | M AS CACO3 | | 228,42 | | | | | 228.42 |

THE IONIC STRENGTH AFTER ACID IS 0.03098 MOLES/LIVER RAU WATER IDNIC STRENGTH WAS 0.02604 HOLE/LITER

THE OSMOTIC PRESSURE IS ESTIMATED TO BE 13.2 PSI AT 25 DEGREES C. II WOULD REQUIRE 367.60 MG/L OF MCL, OR 493.45 MG/L GF H2504, TO GIVE A FH OF 5.60 THIS WATER AS IS, OR FH ADJUSTED WITH HCL, WOULD BE SATURATED WITH CASUA WHEN CONCENTRATED BY A FACTOR OF223.50 If H2SOA IS USED, THE FACTOR IS 10.96

Appendix F

SUMMARY OF TTHMS REGULATIONS

THE MAXIMUM CONTAMINANT LEVEL (MCL)

0.10 mg/l total trihalomethanes.

APPLICABILITY

Community water systems that add disinfectant to the treatment process (ground and surface).

EFFECTIVE DATES

Systems ≥75,000: 2 years after promulgation (29 Nov 1981)

Systems 10,000 to 75,000: 4 years after promulgation (29 Nov 1983)

Systems <10,000: State's discretion

MONITORING REQUIREMENTS

Analyze an annual average of a minimum of 4 samples per quarter per plant taken on same day. Systems using multiple wells drawing raw water from a single aquifer may, with state approval, be considered one treatment plant for determining the required number of samples.

EFFECTIVE DATES

Systems ≥75,000: 1 year after promulgation (29 Nov 1980)

Systems 10-75,000: 3 years after promulgation (29 Nov 1982)

Systems <10,000: State's discretion

SAMPLES LOCATIONS

25% at extreme of distribution system, 75% at location representative of population distribution.

FREQUENCY

For groundwater systems, reduced monitoring may be appropriate for certain systems; states may reduce the requirements through consideration of appropriate data including demonstration by the system that the maximum total trihalomethane potential (MTP) is less than 0.10 mg/l; the minimum frequency would be one sample per year for MTP. For groundwater systems not meeting the above MTP and for surface water systems, states may reduce the monitoring requirements if after 1 year of data collection, TTHMs levels are consistently below 0.10 mg/l; the minimum frequency would be one sample per quarter for TTHMs. The original frequency would be reinstated if the levels exceed 0.10 mg/l or if the treatment or source is modified.

REPORTING REQUIREMENTS TO STATE

Average of each quarterly analysis, within 30 days; until states have adopted the regulations, reporting will be to EPA unless the state requests receipt of data from the public water systems.

REPORTING REQUIREMENTS TO PUBLIC AND STATE

Running annual average of each quarterly sample if it exceeds MCL as prescribed by the public notification provisions.

OTHER REQUIREMENTS

To ensure microbiological quality, state approval of significant modifications in the treatment process to meet MCL.

AGREEMENT REACHED ON TRIHALOMETHANE RULE

The American Water Works Association (AWWA) and the Environmental Protection Agency (EPA) have agreed to settle a lawsuit challenging EPA's standard of 100 parts per billion (0.10~mg/1) for total trihalomethanes (TTHMs) in drinking water.

Under the agreement, EPA will propose a new rule that responds to many of AWWA's concerns. The new rule would set procedures for obtaining and maintaining a variance from the TTHMs standard, and specify control measures a water system may be required to consider.

AWWA said the treatment techniques known as granular activated carbon (GAC) filtration and biologically activated carbon (BAC) filtration for TTHMs control will not be required by EPA. A community would not need to consider those techniques in order to obtain a variance under the proposed rule.

In addition, EPA has clarified its suggestion that a lower standard of 0.010 to 0.025 mg/l is a goal. The agency's original TTHMs proposal left water systems unsure of what control measures eventually would be necessary. In the proposed rule, EPA says it did not mean "to suggest a status for the 0.10 mg/l of short duration."

The proposed new rule lists two groups of technologies for water utilities. The first group is deemed to be widely recognized technologies that are relatively low cost and within the technical capability of most water systems. The second group is considered to be technologies not generally available, but which a system granted a variance could be required to study, and perhaps install, under a compliance schedule.

Group One technologies are: (1) use of chloramines as alternate disinfectant; (2) use of chlorine dioxide as alternate disinfectant; (3) improving existing clarification; (4) alternating the point of chlorination; (5) intermittent or seasonal use of powdered activated carbon and dosages not to exceed 10 mg/l on an annual average basis.

Group Two technologies are: (1) introduction of off-line water storage; (2) aeration where geographically and environmentally appropriate; (3) introduction of clarification where not currently practiced; (4) con sideration of alternative sources of raw water; (5) use of ozone.

EPA or a state exercising enforcement authority could require any Group One item as a condition for granting and maintaining a variance to the TTHM maximums contaminant level of 0.10 mg/l. A water system could challenge such a ruling by demonstrating that the treatment method was not technically appropriate and feasible, or would result in only a marginal reduction of TTHMs for that system.

Group Two technologies could only be required to be installed if EPA or a state demonstrated that for the affected system, the treatment was technically feasible, economically reasonable and would achieve TTHMs reduction at least equal in value to the cost of obtaining the treatment.

AWWA said it believes the proposed rule will minimize potentially disruptive economic and technological effects of the TTHMs standard on water systems. AWWA said it supports adoption of the new rule.

If EPA adopts the proposal as it now stands, AWWA said it will dismiss its pending petition for review of the TTHMs standard, subject to court approval.

The legal action was filed in January 1980. AWWA argued that EPA had not properly determined what control measures were "generally available" and effective for achieving the standard, and that EPA had not properly taken costs into account. AWWA also questioned EPA's stated goal on lowering the standard substantially in the future.

Appendix G

TTHMS ANALYSIS AND ANALYTICAL RESULTS

Table G-1. Diego Garcia's TTHMs Analysis by PWC, Guam.

Samples Taken 12 June 1981

| | | | | | San | ples Take | <u>en 12 Ju</u> | ine 1981 |
|--|------|---|----------------------------------|--------------------------------------|--------------------------------|---|-------------------------------------|--------------------------|
| Sampling Point | Time | Type Sample | Free Cl ₂ Residual | | (ug/1) CHCl ₂ Br | (ug/1) CHC1Br ₂ | (ug/1). CHBr ₃ | (ug/1) TTHM |
| Effluent of Carbon Filter | 0845 | 100 ml w/o* 100 ml w/o 30 ml w/ * 30 ml w/ | 0.0 | 20 17 NT* NT | <1 1 | <1 <1 | < 2 < 2 | 21 19 |
| | | 100 ml blank w/c 60 ml blank w/ | | 17 NT | 1 | <1 | < 2 | 19 |
| Effluent of Reverse Osmosis | 0900 | 100 ml w/o 100 ml w/o 30 ml w/ 30 ml w/ | 0.0 | 17 17 NT NT | ۷ <u>1</u> ا | <1 <1 | ∠2 ∠2 | 18 19 |
| | | 100 ml blank w/c 60 ml blank w/ | ' | 18 NT | 1 | <1 | ۷2 | 20 |
| Effluent of Deionizer | 0915 | 100 ml w/o 100 ml w/o 30 ml w/ | 0.0 | 15 17 NT | <1 1 | <1 <1 | <2 <2 | 16 19 |
| | | 30 ml w/ 100 ml blank w/c 60 ml blank w/ | , | NT 20 NT | 1 | <1 | ۷2 | 22 |
| Lab Faucet of Treated Water | 0715 | 100 ml w/o 100 ml w/o 30 ml w/ 30 ml w/ 100 ml blank w/o 60 ml blank w/ | 3.0 | 117 127 49 78 20 17 | 125 124 44 88 1 | 130 132 52 100 <1 <1 | 30 30 15 15 <2 <2 | 402 413 160 281 |
| Food Services Cold Storage Area, S-Site most remot point south- east | 1040 | 100 ml w/o 100 ml w/o 30 ml w/ 30 ml w/ 100 ml blank w/o 60 ml blank w/ | 2.0 | 332 283 234 239 17 20 | 174 156 141 144 1 | 124 108 112 112 112 <1 <1 | 15 15 15 15 <2 <2 | 645 562 502 510 |
| Officer's Mess Building, most remote point north | 1115 | 100 ml w/o 100 ml w/o 30 ml w/ 30 ml w/ 100 ml blank w/o 60 ml blank w/ | 2.5 | 146 146 83 83 17 18 | 144 147 90 85 1 | 144 148 102 96 <1 <1 | 30 30 22 22 < 2 < 2 | 464 471 297 286 |
| Air Operations Raw Water Tank | 1010 | 100 ml w/o 100 ml w/o 100 ml blank w/o | 0.0 | 16 16 21 | 1 1 1 | <1 <1 <1 | <2 <2 <2 | |
| MAC Air Terminal Restroom | 1000 | 100 ml w/o 100 ml w/o 30 ml w/ 30 ml w/ 100 ml blank w/o 60 ml blank w/o | 1.8 | 156 166 137 122 18 20 | 118 126 115 103 1 | 104 112 112 100 <1 2 | 15 15 15 < 2 < 2 < 2 | 393 419 379 325 |

^{*} w/o - without thiosulfate w/ - with thiosulfate

NT - not tested

Table G-2. TTHMs Analytical Results

[Samples received 9 July 1981; date in ppb]

| | - | (| | | Š | Sample Number | √umb e | ي | | | | | | | | | | |
|--|----|-----------------|-----|-----|----------|---------------|---------------|--------|-----|----------|----------|-----|----------|--------------|-----|-----------|-----|----------|
| 5 37 13 2 25 18 2 2 2 17 14 2 46 22 2 8 <1 | 1_ | - L | 20 | 2.4 | l | 20 | 34 | 38 | 30 | 4A | 48 | 40 | 5A | 28 | 25 | 6A | 89 | 99 |
| 2 17 14 2 46 22 2 8 <1 4 10 <1 33 13 <1 7 <1 <1 3 <1 14 6 <1 2 <1 <1 3 <1 14 6 <1 2 <1 <58 40 4 118 59 4 21 <0 3.0 3.0 0 3.0 0 0 | + | | 13 | 2 | 25 | 18 | 2 | 28 | 12 | 1 | 4 | 5 | 2 | - | - | - | 2 | 7 |
| <1 4 10 <1 33 13 <1 7 <1 <1 3 <1 14 6 <1 2 7 <58 40 4 118 59 4 21 0 3.0 3.0 0 3.0 0 0 | | | 14 | 2 | 46 | 22 | 2 | 84 | 21 | <u>^</u> | 4 | - | ж | ب | | | 4 | - |
| <1 <1 3 <1 14 6 <1 2 7 58 40 4 118 59 4 21 0 3.0 3.0 0 3.0 0 0 | | | 91 | ۲۰ | 33 | 13 | 4 | 74 | 17 | <u>.</u> | <1 | - | 7 | ∵ ∵ | دًا | 41 | Ç | <1 |
| 7 58 40 4 118 59 4 21 0 3.0 3.0 0 3.0 0 0 0 | | | м | | 14 | 9 | 7 | 29 | 6 | 7 | <u>^</u> | دًا | <u>^</u> | ₽ | Ç | ₽ | ₽ | † |
| 0 3.0 3.0 0 3.0 0 | | | 40 | 4 | 118 | 29 | 4 | 215 | 59 | - | 7 | 2 | 6 | 'n | 2 | 2 | 9 | 2 |
| Pappy | _ | | 3.0 | | 3.0 | | 0 | 3.0 | 3.0 | 0 | 3.0 | 3.0 | 0 | 3.0 | 3.0 | 0 | 3.0 | 3.0 |
| No No Yes No No Yes No | | | | Š | S | Yes | <u>Q</u> | O Z | Yes | 2 | 0 | Yes | °Z | 8 | Yes | <u>Q</u> | 2 | Yes |

"B" and "C" sample results are averages of duplicate analyses of the same sample. "A" samples are single samples. Analyses performed 13-15 July 1981. 3 Note:

(2)

Table G-3. TTHMs Analytical Results

[Samples received 22 July, data in ppb]

| D==4===4===4 | | | | | | | | Sar | nple | Num | ber | | | | | | | |
|----------------------|----|----|----|----|----|----|----|-----|------|-----|-----|----|----|----|-------------|----|----|----|
| Pretreatment | 1A | 18 | C4 | 2A | 2B | 2C | 3A | 3B | 3C | 4A | 4B | 4C | 5A | 5B | 5C | 6A | 6B | 60 |
| CHC13 | 1 | 2 | 3 | <1 | 2 | <1 | 3 | 2 | 2 | <1 | 1 | 2 | 3 | 3 | 3 | <1 | 11 | 6 |
| CHC1 ₂ Br | 1 | 1 | 1 | <1 | 1 | 1 | 1 | 2 | 1 | <1 | <1 | <1 | <1 | <1 | <1 | <1 | 1 | <1 |
| CHC1Br ₂ | <1 | <1 | <1 | <1 | <1 | <1 | 3 | <1 | 1 | <1 | <1 | <1 | <1 | <1 | 2 | <1 | <1 | <1 |
| CHBr ₃ | <1 | <1 | <1 | <1 | <1 | <1 | 1 | <1 | <1 | <1 | <1 | <1 | <1 | <1 | <1 | <1 | <1 | <1 |
| Total THM | 2 | 3 | 4 | <1 | 3 | 1 | 8 | 4 | 4 | <1 | 1 | 2 | 3 | 3 | 5 | <1 | 12 | 6 |

Note: Analyses performed 23-25 July 1981. A, B, C, samples were respectively pretreated with 0, 3, 3 ppm of chlorine and no, no, yes of $Na_2S_2O_3$ (addition).

Table G-4. TTHMs Analytical Results

| | | | | | | Sample | es re | [Samples received 9 September 1981, data in ppb] | 9 Sep | tember | 1981, | data | in ppb | | | | | | | |
|--|----|---------------|------|----|------|--------|----------|--|-------|--------|---------------|------|--------|-----|----------|----|--------------------------------|-----|-------|---------------|
| | | | | | | | |) | , | Samp | Sample Number | er | | | | | | | | |
| Pretreatment | 14 | 18 | 10 | 2A | 28 | 2C 3A | | 38 | 30 | 4A | 48* 4C | | 5A | 58 | 25 | 6A | 5C 6A 68** 6C | 99 | Blank | Blank Toluene |
| CHC1 ₃ | - | <1 7.2 6.4 | 6.4 | | 9.2 | 7.2 | 1 | 1.0 9.2 7.2 <1 12.5 6.7 1.0 2.8 2.2 1.0 2.9 2.4 <1 3.9 2.1 | 6.7 | 1.0 | 2.8 | 2.2 | 1.0 | 2.9 | 2.4 | 41 | 3.9 | 2.1 | 41 | ₽ |
| CHC1 ₂ Br | 4 | <1 10.8 6.7 | 6.7 | ٥ | 13.5 | 7.7 | <u>^</u> | 13.5 7.7 <1 46.0 14.0 <1 | 14.0 | | 3.3 | ₽ | ۲ | ₽ | ₽ | ₽ | 3,3 <1 <1 <1 <1 <1 <1 <1 <1 <1 | ۲ | ⊽ | ۲ |
| CHC1Br ₂ | ₽ | <1 6.8 4.1 <1 | 4.1 | ₽ | 7.9 | 4.3 | 7 | 7.9 4.3 <1 48.6 15.0 <1 | 15.0 | | ₽ | ₽ | 4 4 4 | ₽ | Ç | 41 | 4 4 4 4 4 | ₽ | ₽ | ۲ |
| CHBr ₃ | | <1 2.0 1.0 <1 | 1.0 | ⊽ | 2.0 | 1.0 | 7 | 2.0 1.0 <1 15.0 6.5 <1 | 6.5 | ۵. | 4.5 <1 <1 | ₽ | ۵ | ₽ | <u>^</u> | 41 | <1 <1 <1 4.9 <1 | û | ₽ | ۵. |
| Total TTHMs <1 26.8 18.2 | ₽ | 26.8 | 18.2 | | 32.6 | 20.2 | ₽ | 1.0 32.6 20.2 <1 121.8 42.2 1.0 10.6 2.2 1.0 2.9 2.4 <1 11.1 2.1 | 42.2 | 1.0 | 10.6 | 2.2 | 1.0 | 5.9 | 2.4 | 41 | 11.1 | 2.1 | | |
| Cl ₂ added, ² ppm | 0 | 4 | 4 | 0 | 4 | 4 | 0 | 4 4 | 4 | 0 | 4 | | 0 | 4 | 4 | 0 | 4 | 4 | | |

Note: * Set 4B samples-11 unidentified peaks. **Set 6B-10 unidentified peaks.

Appendix H

ACIDS FOR ph CONTROL IN RO PROCESS

Two acids readily available in commercial quantities (55-gallon drums or bulk) are hydrochloric acid and sulfuric acid. Hydrochloric acid is usually sold as 20° Baumé or about 31.4% HCl. Sulfuric acid can be obtained in several different concentrations, the most common being battery electrolyte (33.3%), concentrated (94%), and oleum (100%).

In calculating the amount of each acid required to neutralize a given alkalinity, the following chemical reactions apply:

$$2(36.5)$$
 162
 $2HC1 + Ca(HCO_3)_2 = 2H_2O + CaCl_2 + 2CO_2$ (1)
 0.45 lb 1 lb

$$\begin{array}{rcl}
98 & 162 \\
H_2SO_4 + Ca(HCO_3)_2 &= 2H_2O + CaSO_4 + 2CO_2 \\
0.6 lb & 1 lb
\end{array} \tag{2}$$

The specific gravity of 20°Be HCl is 1.16 and that of concentrated H_2SO_4 is 1.83 or 9.6 lb/gal for HCl and 15.2 lb/gal for H_2SO_4 .
Therefore, it would require 0.15 gallons of HCl (31.4%) and/or

0.042 gallons of ${\rm H_2SO_A}$ (94%) to neutralize 1 pound alkalinity as

Ca(HCO₃)₂. For a 450,000-gallon per day water treatment system, assuming alkalinity concentration of 500 ppm as $CaCO_3$ (equivalent to 810 ppm as $Ca(HCO_3)_2$, the total alkalinity content is 3,025 pounds. This alkalinity will require 454 gallons of HCl (31.4%) and/or 127 gallons of H₂SO_A (94%) to neutralize. The HCl requirement found in the actual ffeld test was 15 gallons HC1/26,000 gallons of feed water or 0.057% (v/v) that was equivalent to 0.081% (v/v) for product water at 70% recovery rate. Based on this figure, 366 gallons of 20°Be HCl will be needed for obtaining 450,000 gallons of product water per day at a 70% recovery rate. The difference between 454 and 366 gallons is a result of the assumption that alkalinity is all in $Ca(HCO_2)_2$ form rather than in many other forms.

Appendix I

LANGLIER'S SATURATION INDEX (LI)

$$LI = pH - pH_{e}$$
 (1)

$$pH_s = pK_2 - pK_s + p[Ca] + p[A]k] + 6.301 + S$$
 (2)

$$S = 2.5 (\mu)^{0.5}/[1 + 5.3 (\mu)^{0.5} + 5.5\mu]$$
 (3)

pH = pH value of the water sample

where $pH_s = pH$ of saturation

p = common logarithm

 K_2 = the second ionization constant for carbonic acid

 K_e = the solubility product constant

[Ca] = calcium hardness in ppm as $CaCO_2/50$

[Alk] = alkalinity in ppm as $CaCO_3/50$

 $S = 2.5 \times 10^{-5} \text{Sd}$; Sd = TDS in ppm (when less than 500)

 $S \approx 4H-A \text{ when } Sd > 500 \text{ ppm}(TDS)$

H = total hardness in moles/1

A = alkalinity in eq/1

The carbonate solution equilibrium constants vary with temperature as follows:

$$Log (K_s/K_2)$$
 +2.47 +2.34 +2.21 +2.10 +1.99 +1.71 +1.40

A linear regression of these values can be expressed as:

$$Log (K_s/K_2) = 2.55 - 0.0216T$$
 (4)

There is no corrosion and scaling when LI=0: LI with a positive value indicates scaling tendency, while LI with a negative value indicates corrosion tendency.

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